


04/27/98

Jc135 U.S. PTO

A

TRANSMITTAL OF UTILITY APPLICATION UNDER 37 C.F.R. §1.53	Attorney Docket No.	7466-2301
	First named inventor	Hubert Köster
	Express mail label #	EM468592435US
	Date of mailing	April 27, 1998

Application Elements	Accompanying Application Papers
1. <input checked="" type="checkbox"/> Fee Transmittal Form 2. <input checked="" type="checkbox"/> Specification No. Pages <u>78</u> (including Abstract) a. Title: SOLUTION PHASE BIOPOLYMER SYNTHESIS b. Number of claims: <u>49</u> 3. <input type="checkbox"/> No. sheets of drawings <u>0</u> with <u>0</u> Figs. 4. <input checked="" type="checkbox"/> Unexecuted Declaration listing names of joint inventors 5. <input type="checkbox"/> Sequence Listing <input type="checkbox"/> Paper copy (identical to computer copy) <input type="checkbox"/> Computer readable copy <input type="checkbox"/> Verified statement	6. <input type="checkbox"/> Assignment papers 7. <input type="checkbox"/> Statement of status as small entity 8. <input checked="" type="checkbox"/> Return Receipt Postcard
	SIGNATURE OF ATTORNEY/AGENT BROWN MARTIN HALLER & McCLAIN  Stephanie Seidman Registration Number: 33,779
If a continuing application: N/A <input type="checkbox"/> continuation <input type="checkbox"/> Divisional <input type="checkbox"/> continuation-in-part of prior U.S. application Serial No.	
CORRESPONDENCE ADDRESS	
NAME	Stephanie Seidman Registration No. 33,779 Brown, Martin, Haller & McClain
Address	1660 Union Street, San Diego, California 92101
	Telephone: 619/238-0999 Facsimile: 619/238-0062

09053227 00423700

FEE TRANSMITTAL ACCOMPANYING UTILITY APPLICATION UNDER 37 C.F.R. §1.53	Attorney Docket No.	7466-2301
	First named inventor	Hubert Köster
	Express mail label #	EM468592435US
	Date of mailing	April 27, 1998

FEE CALCULATION FOR CLAIMS AS FILED

a)	Basic Fee		\$ 790.00
b)	Independent Claims <u>3</u> - 3 = <u>0</u> x \$ 82.00		\$.00
c)	Total Claims <u>49</u> - 20 = <u>29</u> x \$ 22.00		\$ 638.00
d)	Fee for Multiple Dependent Claims - \$260.00		\$ 0.00
TOTAL FILING FEE			\$ 1428.00

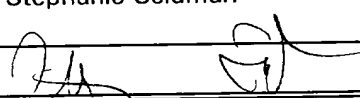
[X] Statement(s) of Status as Small Entity
reducing Filing Fee by one-half to
(to be submitted under separate cover) \$714.00

[X] A check in the amount of \$714.00 to cover the fee for filing the application.

[] Charge \$____ to Deposit Account No. 02-4070.

[X] The Commissioner is hereby authorized to charge any fees, including the filing fee and additional claim fees, that may be required in this application under 37 C.F.R. §§ 1.16-1.17 during its entire pendency, or credit any overpayment, to Deposit Account No. 02-4070. If proper payment is not enclosed, such as a check in the wrong amount, unsigned, post-dated, otherwise improper or informal, or absent, the Commissioner is authorized to charge the unpaid amount to Deposit Account No. 02-4070 during the entire pendency of this application. This sheet is filed in duplicate.

CORRESPONDENCE ADDRESS	
NAME	Stephanie Seidman Registration No. 33,779 Brown, Martin, Haller & McClain
Address	1660 Union Street, San Diego, California 92101
	Telephone: 619/238-0999 Facsimile: 619/238-0062

Submitted by:			
Typed or printed name	Stephanie Seidman	Reg. Number	33,779
Signature		Date	04/27/98
		Deposit Account	02-4070

04/27/98 14:22:00

SOLUTION PHASE BIOPOLYMER SYNTHESIS

FIELD OF INVENTION

Liquid phase carriers (LPCs) and methods of use of LPCs in biopolymer synthesis are provided. In particular, multifunctional LPCs and
 5 methods of use of multifunctional LPCs in oligonucleotide synthesis are provided.

BACKGROUND OF THE INVENTION

Recent developments in chemistry and biology are indicative of an increased demand for modified biopolymers, such as oligonucleotides,
 10 peptides and oligosaccharides. For most applications, oligonucleotides are needed only in small amounts (nano- to milligrams), for use in, e.g., polymerase chain reactions (see, e.g., K.B. Mullis Angew. Chem. **1994**, 106, 1271-1276 and K.B. Mullis et al. Methods Enzymol. **1987**, 155, 335-3508) and as diagnostic tools (see, e.g., S. Beck et al. Anal. Chem.
 15 **1990**, 62, 2258-2270; C. Jurinke et al. Anal. Biochem. **1996**, 237, 174-181; C. Jurinke et al. Genetic Analysis: Biomolecular Engineering **1996**, 13, 67-71; C. Jurinke et al. Anal. Biochem. **1997**, 69, 904-910 and C.W. Siegert et al. Anal. Biochem. **1996**, 243, 55-65).

There, however, is an increase in uses of biopolymers as
 20 therapeutic agents. For example, antisense oligonucleotides (see, e.g., E. Uhlmann et al. Chem. Rev. **1990**, 90, 543-584 and F.F. Milligan et al. J. Med. Chem. **1993**, 36, 1923-1937) may be used as therapeutic agents. Current methods of biopolymer synthesis, such as solid phase synthesis of oligonucleotides and peptides, result in preparation of small amounts of
 25 a desired biopolymer (see, e.g., N. D. Sinha et al. Tetrahedron Lett. **1983**, 24, 5843-5846; H. Köster et al. Tetrahedron **1984**, 40, 103-112; and N.D. Sinha et al. Nucleic Acids Res. **1984**, 12, 4539-4557).

Thus, there is a need for economical large-scale methods of synthesis of modified biopolymers, such as oligonucleotides, peptides and
 30 oligosaccharides, that provide the biopolymer with the requisite high purity for therapeutic use. Synthesis in solution offers advantages in

large scale production, such as easy upscaling, direct reaction control and purification after each reaction cycle, affording products of high purity.

One drawback of solution phase synthesis lies in lower yields, caused by loss of product during purification and incomplete reactions, because

- 5 reagents cannot be used in large excesses. Known methods of solution phase synthesis of biopolymers also suffer from loss of yield as the length of the biopolymeric chain increased.

- A further drawback of solution phase techniques is the lack of solubility of the substrate biopolymer in organic solvents when the
10 number of monomeric units exceeds 3-5. Such lack of solubility also contributes to lower yields seen in solution phase synthesis of biopolymers. Therefore, there is a need for carriers that mediate the solubility of the growing biopolymeric chain.

- Therefore, it is an object herein to provide carriers for solution
15 phase synthesis of biopolymers. It is also an object herein to provide methods for biopolymer synthesis using the carriers.

SUMMARY OF THE INVENTION

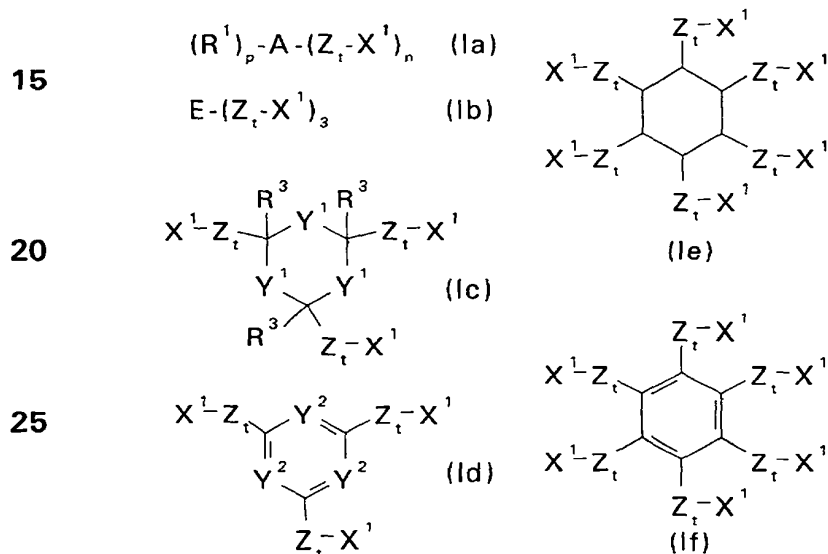
- Liquid phase carriers (LPCs), LPCs coupled to biopolymers and methods of synthesis of biopolymers using LPCs are provided. In
20 particular, biopolymers that can be prepared using the LPCs and methods described herein include, but are not limited to, oligonucleotides, peptides, peptide nucleic acids (PNAs) and oligosaccharides. A desired biopolymer may be prepared by the methods provided herein, or modification thereof using techniques known to those of skill in the art.

- 25 The LPCs provided herein are multifunctional compounds that allow for the simultaneous construction of three or more biopolymers, which may be of differing structure, but more typically are identical in structure.

- The LPCs have the formula $Sp(X^1)_n$. Sp refers to a group that serves as the scaffold upon which synthesis of a plurality of oligomers is
30 effected. X_1 is the reactive group for effecting synthesis of a selected biopolymer, and "n" is preferably 3 or more. In particular, Sp is a

polyvalent group, such as an atomic group, a cyclic group or an aromatic group (heterocycles, carbocycles, aryls, heteroaryls, such that the resulting structure is symmetrically disposed around the center of the cyclic group), that has more than two points of attachment, n is the number of points of attachment in Sp, and is preferably 3-6, and X¹ is a reactive group for performing synthesis of a biopolymer. Sp is preferably a symmetrical group such that all X¹ groups are equivalent. It will be appreciated by those of skill in the art that such symmetry allows for uniformity in the synthesis of n biopolymers, where n is preferably 3 or more, using the LPCs provided herein. The LPCs provided herein are intended for use for synthesis of three or more biopolymers.

In one embodiment, the LPCs have one of formulae (I):



30 where Sp is (R¹)_p-A, E or a cyclic group (i.e. heterocycles, carbocycles, aryls, heteroaryls, such that the resulting structure is symmetrically disposed around the center of the cyclic group) with the linked "Z_t" moieties, where t is 0 or 1. As depicted the cyclic and aromatic rings

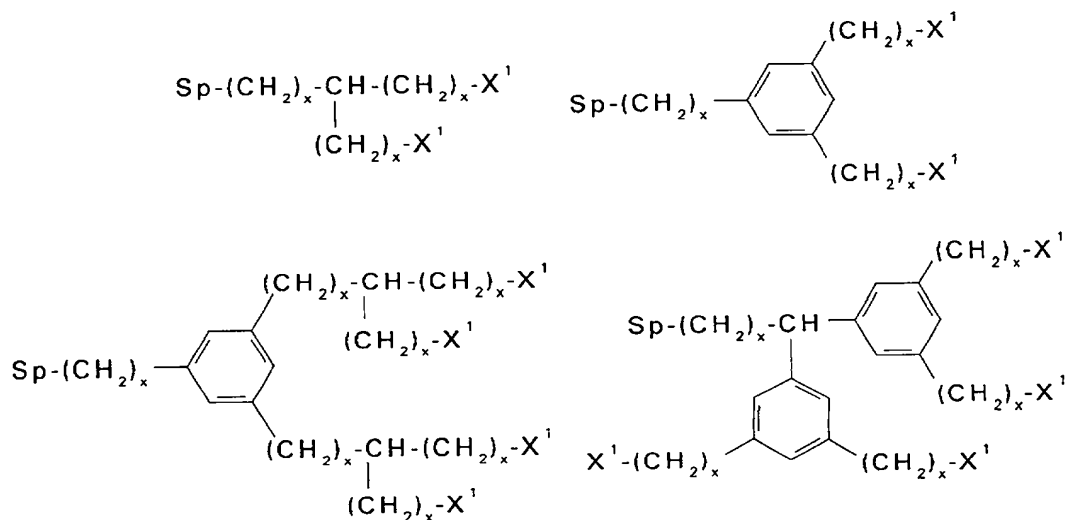
35 have 6 members; it is understood that fewer or higher-number membered rings may also be used, as long as the resulting structure possesses the

requisite symmetry, and the number of linkages for synthesis of the biopolymers is greater than two.

- In particular, A is carbon or silicon; E is nitrogen or P(O); R¹ and R³ are each independently hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocyclalkyl; p is 0 or 1; Z, where t is 0 or 1, is a divalent hydrocarbon, containing one or more or mixtures of phenylene or alkylene groups, and contains a sufficient number of carbons (0, where t is 0, up to as many as 24 or more) to prevent or to reduce interactions among the chains in each biopolymer during synthesis. Preferably Z, typically containing from 1 up to 30 carbons or more, is any combination alkylene and arylene units, preferably methylene and phenylene units, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12 units, preferably selected from 1,2-, 1,3- or 1,4-, preferably 1,4-, phenylene and alkylene units, more preferably methylene units, which units may be combined in any order; X¹ is any reactive group that is used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably halide, OH, SH, NH₂, COR⁵ or COOR⁴; n is preferably 3 or 4; R⁴ is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclalkyl; R⁵ is halide, heteroaryl or pseudohalide; Y¹ is CH₂, NH, S or O; and Y² is selected from CH and N.

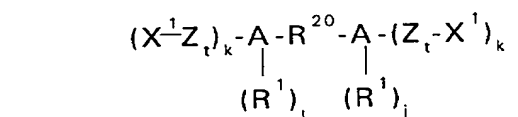
- In other embodiments, Z is a multivalent group, particularly a trivalent or higher valency hydrocarbon, with three or more points of attachment: The points of attachment are to A, E, or the cyclic nucleus, and the others are to two or more X¹ groups. In these embodiments, Z is chosen such that the symmetry of the LPC is maintained and all of the X¹ groups are equivalent.

- Thus, the resulting LPCs, in these embodiments, are chosen from compounds such as, but not limited to, the following:

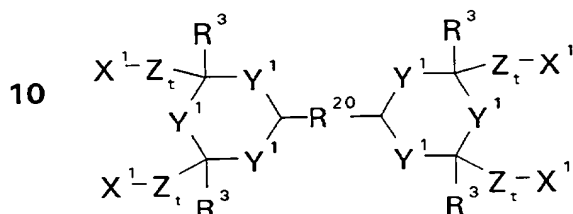
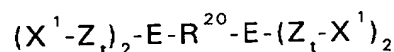


where Sp (*i.e.*, $(R^1)_p$ -A, E or the cyclic group with the linked " Z_t " moieties, where t is 0 or 1) and X^1 are chosen as above and x is preferably 0-12, more preferably 0-6.

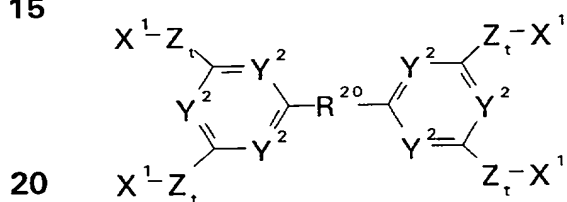
- In certain embodiments, the LPCs contain two or more of the
- 5 compounds of formula (I) that are linked together so that the symmetry of the resulting LPC is maintained (*i.e.*, all X^1 groups are equivalent). In preferred embodiments, the LPCs have formulae:



5



15



20

where A, E, R¹, R³, X¹, Y¹, Y², t and Z are defined as for formula (I); R²⁰ is
 alkylene, alkenylene, alkynylene, arylene or heteroarylene of any desired
 length, preferably from 0 to 24, more preferably from 1 to about 12; k is
 2 or 3; and j is 0 or 1.

Any of the groups R¹, R³, R⁴, R⁵, X¹, Y¹, Y² and Z may be
 unsubstituted or substituted with one or more substituents each inde-
 pendently selected from Q, which, as defined herein, is halogen, hydroxy,
 nitrile, nitro, formyl, mercapto, carboxy, alkyl, haloalkyl, polyhaloalkyl,
 aminoalkyl, diaminoalkyl, alkenyl containing 1 to 2 double bonds, alkynyl
 containing 1 to 2 triple bonds, cycloalkyl, cycloalkylalkyl, aryl, heteroaryl,
 arylalkyl, heteroarylalkyl, alkylidene, arylalkylidene, alkylcarbonyl,
 arylcarbonyl, heteroarylcarbonyl, alkoxycarbonyl, alkoxycarbonylalkyl,
 aryloxycarbonyl, aryloxycarbonylalkyl, aminocarbonyl, alkylaminocar-
 bonyl, dialkylaminocarbonyl, arylaminocarbonyl, diarylaminocarbonyl, aryl-
 alkylaminocarbonyl, alkoxy, aryloxy, perfluoroalkoxy, alkenyloxy, alkynyl-
 oxy, arylalkoxy, amino, aminoalkyl, alkylaminoalkyl, dialkylaminoalkyl,

arylaminoalkyl, diarylaminoalkyl, alkylamino, dialkylamino, arylamino, diarylamino, alkylarylamino, alkylcarbonylamino, alkoxycarbonylamino, arylcarbonylamino, aryloxycarbonylamino, azido, alkylthio, arylthio, perfluoroalkylthio, thiocyano, isothiocyano, alkylsulfinyl, alkylsulfonyl, 5 arylsulfinyl, arylsulfonyl, aminosulfonyl, alkylaminosulfonyl, dialkylamino-sulfonyl, arylaminosulfonyl or diarylamino-sulfonyl.

In certain embodiments, the LPCs are chosen as above, with the proviso that Z contains more than two methylene groups. In other 10 embodiments, the LPCs are chosen with the proviso that Z is not 1,4-arylene. In more preferred embodiments, the LPCs are chosen with the proviso that Z contains more than two methylene groups and is not 1,4-arylene. In other more preferred embodiments, the LPCs are chosen with the proviso that Z contains at least one, preferably at least two, more preferably at least three or more phenylene or alkylene units.

15 Methods of synthesis of biopolymers using the LPCs described herein are also provided. The LPCs and methods permit synthesis of relatively large amounts, i.e. therapeutically useful amounts or preparative amounts (on the order of at least 0.1 grams, preferably a gram or more in volumes at which solid phase methods and other liquid phase methods 20 produce substantially lower amounts) of biopolymers, including, but not limited to, oligonucleotides, peptides, peptide nucleic acids (PNAs) and oligosaccharides, may be produced by the methods. It is possible to prepare not only naturally occurring biopolymers, but also analogs of the biopolymers. Such analogs include, for example, analogs of oligonucleo- 25 tides such as oligo-triesters or oligophosphonates. Use of the LPCs provided herein in the methods afford biopolymers of high purity and in significantly higher yields than methods known in the art.

In one embodiment, the methods involve the steps of (a) reacting an LPC of formula $Sp(X^1)_n$, provided herein, with a first monomer N^1 ; (b) 30 separating and purifying the product of step (a) to afford a compound of formula $Sp(X^1-N^1)_n$; (c) reacting the product of step (b) with a second

- monomer N^2 , a dimer N^2-N^3 or a trimer $N^2-N^3-N^4$; and (d) repeating steps (b) and (c) using the desired monomers, dimers or trimers to produce the desired LPC-bound biopolymer of formula $Sp(X^1-N^1-N^2-...-N^m)_n$, where m is 3 to 100, preferably 3-50, more preferably 3-25, most preferably 3-10. In
- 5 another embodiment, the methods further involve the step of (e) cleaving the biopolymer from the LPC.

In another embodiment, the methods involve the steps of:

- (a) reacting an LPC of formula $Sp(X^1)_n$ with a first monomer N^1 ;
- (b) separating and purifying the product of step (a) to afford a
- 10 compound of formula $Sp(X^1-N^1)_n$;
- (c) reacting the product of step (b) with a second monomer N^2 , a dimer N^2-N^3 or a trimer $N^2-N^3-N^4$; and
- (d) repeating steps (b) and (c) to produce an LPC-bound biopolymer, preferably an oligonucleotide, of formula $Sp(X^1-N^1-N^2-...-N^m)_n$,
- 15 where m is 3 to 100, wherein:

Sp is a polyvalent group that has two or more points of attachment, n corresponds to the number of points of attachment in Sp and X^1 is a reactive group for biopolymer, preferably oligonucleotide, synthesis;

- 20 $N^1, N^2, N^3...N^m$ are biopolymer monomers, preferably nucleotides or nucleosides;

the dimers and trimers comprise the monomers; and

the protocol used in steps (c) and (d) to synthesize the biopolymer, preferably the oligonucleotide, is the phosphoramidite protocol.

- 25 Monomers (N) used in preparing the biopolymer will depend on the biopolymer desired. For example, monomers which are used in the methods include nucleotides, nucleosides, natural and unnatural amino acids, PNA monomers and monosaccharides. In the methods, synthesis of the biopolymeric chain (i.e., steps c and d) may be performed by any
- 30 protocol known to those of skill in the art. A preferred protocol for oligonucleotide synthesis is the phosphoramidite protocol (see, e.g., S. L.

Beaucage *et al.* *Tetrahedron* **1992**, *48*, 2223-2311 and S. L. Beaucage *et al.* *Tetrahedron Lett.* **1981**, *22*, 1859-1862).

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Definitions

5 Unless defined otherwise, all technical and scientific terms used herein have the same meaning as is commonly understood by one of skill in the art to which this invention belongs. All patents and publications referred to herein are incorporated by reference.

As used herein, peptide nucleic acid (PNA) refers to nucleic acid
10 analogs in which the ribose-phosphate backbone is replaced by a backbone held together by amide bonds.

As used herein, MALDI-TOF refers to matrix assisted laser desorption ionization-time of flight mass spectrometry.

As used herein, a liquid phase carrier (LPC) is a symmetrical
15 compound that possesses functionality compatible with biopolymer synthesis. Such functionality includes, but is not limited to, alcohol, thiol, amino, carboxylic acid, acyl halide, ester, acyl heteroaryl and acyl pseudohalide groups. Preferably, the symmetry of the LPC is such that all of the functional groups are equivalent. Multifunctional LPCs are LPCs
20 that possess three or more functional groups for use in biopolymer synthesis.

As used herein, a biopolymer is any compound found in nature, or derivatives thereof, made up of monomeric units. Biopolymers include, but are not limited to, oligonucleotides, peptides, peptide nucleic acids
25 (PNAs) and oligosaccharides. Thus, the monomeric units include, but are not limited to, nucleotides, nucleosides, amino acids, PNA monomers, monosaccharides, and derivatives thereof.

As used herein, substantially pure means sufficiently homogeneous to appear free of readily detectable impurities as determined by standard
30 methods of analysis, such as thin layer chromatography (TLC), gel electrophoresis, high performance liquid chromatography (HPLC) and

mass spectrometry (MS), used by those of skill in the art to assess such purity, or sufficiently pure such that further purification would not detectably alter the physical and chemical properties, such as enzymatic and biological activities, of the substance. Methods for purification of the compounds to produce substantially chemically pure compounds are known to those of skill in the art. A substantially chemically pure compound may, however, be a mixture of stereoisomers. In such instances, further purification might increase the specific activity of the compound.

As used herein, alkyl, alkenyl and alkynyl carbon chains, if not specified contain from 1 to 20 carbons, preferably 1 to 16 carbons, and are straight or branched. Alkenyl carbon chains of from 2 to 20 carbons preferably contain 1 to 8 double bonds, and the alkenyl carbon chains of 1 to 16 carbons preferably contain 1 to 5 double bonds. Alkynyl carbon chains of from 2 to 20 carbons preferably contain 1 to 8 triple bonds, and the alkynyl carbon chains of 2 to 16 carbons preferably contain 1 to 5 triple bonds. The alkyl, alkenyl and alkynyl groups may be optionally substituted, with one or more groups, preferably alkyl group substituents that may be the same or different. As used herein, lower alkyl, lower alkenyl, and lower alkynyl refer to carbon chains having less than about 6 carbons.

As used herein, an "alkyl group substituent" includes halo, haloalkyl, preferably halo lower alkyl, aryl, hydroxy, alkoxy, aryloxy, alkyloxy, alkylthio, arylthio, aralkyloxy, aralkylthio, carboxy alkoxy, carbonyl, oxo and cycloalkyl.

As used herein, "aryl" refers to cyclic groups containing from 3 to 19 carbon atoms. Aryl groups include, but are not limited to groups, such as phenyl, substituted phenyl, naphthyl, substituted naphthyl, in which the substituent is lower alkyl, halogen, or lower alkoxy.

As used herein, an "aryl group substituent" includes alkyl, cycloalkyl, cycloalkylalkyl, aryl, heteroaryl optionally substituted with 1 or

- more, preferably 1 to 3, substituents selected from halo, halo alkyl and alkyl, arylalkyl, heteroarylalkyl, alkenyl containing 1 to 2 double bonds, alkynyl containing 1 to 2 triple bonds, halo, hydroxy, haloalkyl and polyhaloalkyl, preferably halo lower alkyl, especially trifluoromethyl,
- 5 formyl, alkylcarbonyl, arylcarbonyl that is optionally substituted with 1 or more, preferably 1 to 3, substituents selected from halo, halo alkyl and alkyl, heteroarylcarbonyl, carboxy, alkoxycarbonyl, aryloxy, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, arylamino-
- 10 carbonyl, diarylaminocarbonyl, arylalkylaminocarbonyl, alkoxy, aryloxy, perfluoroalkoxy, alkenyloxy, alkynyloxy, arylalkoxy, aminoalkyl, alkyl-
- aminoalkyl, dialkylaminoalkyl, arylaminoalkyl, amino, alkylamino, dialkyl-
- amino, arylamino, alkylaryl amino, alkylcarbonylamino, arylcarbonylamino, azido, nitro, mercapto, alkylthio, arylthio, perfluoroalkylthio, thiocyno, isothiocyno, alkylsulfinyl, alkylsulfonyl, arylsulfinyl, arylsulfonyl,
- 15 aminosulfonyl, alkylaminosulfonyl, dialkylaminosulfonyl and arylamino-
- sulfonyl.

- As used herein, "cycloalkyl" refers to a saturated mono- or multi-
- cyclic ring system, preferably of 3 to 10 carbon atoms, more preferably 3
- 20 to 6 carbon atoms; cycloalkenyl and cycloalkynyl refer to mono- or
- multicyclic ring systems that respectively include at least one double bond
- and at least one triple bond. Cycloalkenyl and cycloalkynyl groups may
- preferably contain 3 to 10 carbon atoms, with cycloalkenyl groups more
- preferably containing 4 to 7 carbon atoms and cycloalkynyl groups more
- preferably containing 8 to 10 carbon atoms. The ring systems of the
- 25 cycloalkyl, cycloalkenyl and cycloalkynyl groups may be composed of one
- ring or two or more rings which may be joined together in a fused,
- bridged or spiro-connected fashion, and may be optionally substituted
- with one or more alkyl group substituents.

- As used herein, "heteroaryl" refers to a monocyclic or multicyclic
- 30 ring system, preferably of about 5 to about 15 members where one or
- more, more preferably 1 to 3 of the atoms in the ring system is a

heteroatom, that is, an element other than carbon, for example, nitrogen, oxygen and sulfur atoms. The heteroaryl may be optionally substituted with one or more, preferably 1 to 3, aryl group substituents. Exemplary heteroaryl groups include, for example, furyl, imidazynyl, pyrrolidinyl, pyrimidinyl, tetrazolyl, thienyl, pyridyl, pyrrolyl, N-methylpyrrolyl, quinolinyl and isoquinolinyl, with pyridyl and quinolinyl being preferred.

As used herein, "heterocyclic" refers to a monocyclic or multicyclic ring system, preferably of 3 to 10 members, more preferably 4 to 7 members, even more preferably 5 to 6 members, where one or more, preferably 1 to 3 of the atoms in the ring system is a heteroatom, that is, an element other than carbon, for example, nitrogen, oxygen and sulfur atoms. The heterocycle may be optionally substituted with one or more, preferably 1 to 3 aryl group substituents. Preferred substituents of the heterocyclic group include hydroxy, amino, alkoxy containing 1 to 4 carbon atoms, halo lower alkyl, including trihalomethyl, such as trifluoromethyl, and halogen. As used herein, the term heterocycle may include reference to heteroaryl.

As used herein, the nomenclature alkyl, alkoxy, carbonyl, etc. are used as is generally understood by those of skill in this art. For example, as used herein alkyl refers to saturated carbon chains that contain one or more carbons; the chains may be straight or branched or include cyclic portions or be cyclic. As used herein, alicyclic refers to aryl groups that are cyclic.

As used herein, "halogen" or "halide" refers to F, Cl, Br or I. As used herein, pseudohalides are compounds that behave substantially similar to halides. Such compounds can be used in the same manner and treated in the same manner as halides (X^- , in which X is a halogen, such as Cl or Br). Pseudohalides include, but are not limited to cyanide, cyanate, thiocyanate, selenocyanate, trifluoromethyl and azide.

As used herein, "haloalkyl" refers to a lower alkyl radical in which one or more of the hydrogen atoms are replaced by halogen including, but

not limited to, chloromethyl, trifluoromethyl, 1-chloro-2-fluoroethyl and the like.

As used herein, "haloalkoxy" refers to RO- in which R is a haloalkyl group.

- 5 As used herein, "sulfinyl" refers to -S(O)-. As used herein, "sulfonyl" refers to -S(O)₂-.

As used herein, "aminocarbonyl" refers to -C(O)NH₂.

- As used herein, "alkylaminocarbonyl" refers to -C(O)NHR in which R is hydrogen or alkyl, preferably lower alkyl. As used herein "dialkyl-
10 aminocarbonyl" as used herein refers to -C(O)NR'R in which R' and R are independently selected from hydrogen or alkyl, preferably lower alkyl; "carboxamide" refers to groups of formula -NR'COR.

- As used herein, "diarylamino carbonyl" refers to -C(O)NRR' in which R and R' are independently selected from aryl, preferably lower aryl, more preferably phenyl.
15

As used herein, "arylalkylaminocarbonyl" refers to -C(O)NRR' in which one of R and R' is aryl, preferably lower aryl, more preferably phenyl, and the other of R and R' is alkyl, preferably lower alkyl.

- As used herein, "arylamino carbonyl" refers to -C(O)NHR in which R
20 is aryl, preferably lower aryl, more preferably phenyl.

As used herein, "alkoxycarbonyl" refers to -C(O)OR in which R is alkyl, preferably lower alkyl.

As used herein, "aryloxycarbonyl" refers to -C(O)OR in which R is aryl, preferably lower aryl, more preferably phenyl.

- 25 As used herein, "alkoxy" and "alkylthio" refer to RO- and RS-, in which R is alkyl, preferably lower alkyl.

As used herein, "aryloxy" and "arylthio" refer to RO- and RS-, in which R is aryl, preferably lower aryl, more preferably phenyl.

- As used herein, "alkylene" refers to a straight, branched or cyclic,
30 preferably straight or branched, bivalent aliphatic hydrocarbon group, preferably having from 1 to about 20 carbon atoms, more preferably 1 to

- 12 carbons, even more preferably lower alkylene. The alkylene group is optionally substituted with one or more "alkyl group substituents." There may be optionally inserted along the alkylene group one or more oxygen, sulphur or substituted or unsubstituted nitrogen atoms, where the
- 5 nitrogen substituent is alkyl as previously described. Exemplary alkylene groups include methylene ($-\text{CH}_2-$), ethylene ($-\text{CH}_2\text{CH}_2-$), propylene ($-(\text{CH}_2)_3-$), cyclohexylene ($-\text{C}_6\text{H}_{10}-$), methylenedioxy ($-\text{O}-\text{CH}_2-\text{O}-$) and ethylenedioxy ($-\text{O}-(\text{CH}_2)_2-\text{O}-$). The term "lower alkylene" refers to alkylene groups having 1 to 6 carbons. Preferred alkylene groups are lower
- 10 alkylene, with alkylene of 1 to 3 carbon atoms being particularly preferred.

- As used herein, "alkenylene" refers to a straight, branched or cyclic, preferably straight or branched, bivalent aliphatic hydrocarbon group, preferably having from 2 to about 20 carbon atoms and at least
- 15 one double bond, more preferably 1 to 12 carbons, even more preferably lower alkenylene. The alkenylene group is optionally substituted with one or more "alkyl group substituents." There may be optionally inserted along the alkenylene group one or more oxygen, sulphur or substituted or unsubstituted nitrogen atoms, where the nitrogen substituent is alkyl as
- 20 previously described. Exemplary alkenylene groups include $-\text{CH}=\text{CH}-\text{CH}=\text{CH}-$ and $-\text{CH}=\text{CH}-\text{CH}_2-$. The term "lower alkenylene" refers to alkenylene groups having 2 to 6 carbons. Preferred alkenylene groups are lower alkenylene, with alkenylene of 3 to 4 carbon atoms being particularly preferred.

- 25 As used herein, "alkynylene" refers to a straight, branched or cyclic, preferably straight or branched, bivalent aliphatic hydrocarbon group, preferably having from 2 to about 20 carbon atoms and at least one triple bond, more preferably 1 to 12 carbons, even more preferably lower alkynylene. The alkynylene group is optionally substituted with one
- 30 or more "alkyl group substituents." There may be optionally inserted along the alkynylene group one or more oxygen, sulphur or substituted or

unsubstituted nitrogen atoms, where the nitrogen substituent is alkyl as previously described. Exemplary alkynylene groups include

—C≡C—C≡C—, —C≡C— and —C≡C—CH₂—. The term "lower alkynylene"

refers to alkynylene groups having 2 to 6 carbons. Preferred alkynylene groups are lower alkynylene, with alkynylene of 3 to 4 carbon atoms being particularly preferred.

As used herein, "arylene" refers to a monocyclic or polycyclic, preferably monocyclic, bivalent aromatic group, preferably having from 3 to about 20 carbon atoms and at least one aromatic ring, more preferably 3 to 12 carbons, even more preferably lower arylene. The arylene group is optionally substituted with one or more "alkyl group substituents." There may be optionally inserted around the arylene group one or more oxygen, sulphur or substituted or unsubstituted nitrogen atoms, where the nitrogen substituent is alkyl as previously described. Exemplary arylene groups include 1,2-, 1,3- and 1,4-phenylene. The term "lower arylene" refers to arylene groups having 5 or 6 carbons. Preferred arylene groups are lower arylene.

As used herein, "heteroarylene" refers to a bivalent monocyclic or multicyclic ring system, preferably of about 5 to about 15 members where one or more, more preferably 1 to 3 of the atoms in the ring system is a heteroatom, that is, an element other than carbon, for example, nitrogen, oxygen and sulfur atoms. The heteroarylene group may be optionally substituted with one or more, preferably 1 to 3, aryl group substituents.

As used herein, "alkylidene" refers to a bivalent group, such as =CR'R", which is attached to one atom of another group, forming a double bond. Exemplary alkylidene groups are methylidene (=CH₂) and ethylidene (=CHCH₃). As used herein, "arylalkylidene" refers to an alkylidene group in which either R' or R" is an aryl group.

As used herein, "amido" refers to a bivalent group, either —C(O)NH— or —HNC(O)—. "Thioamido" refers to a bivalent group, either —C(S)CH— or —

HNC(S)-. "Oxyamido" refers to a bivalent group, either -OC(O)NH- or -HNC(O)O-. "Thiaamido" refers to a bivalent group, either -SC(O)NH- or -HNC(O)S-. "Dithiaamido" refers to a bivalent group, either -SC(S)NH- or -HNC(S)S-. "Ureido" refers to the bivalent group -HNCONH-.

- 5 "Thioureido" refers to the bivalent group -HNCSNH-.

As used herein, the term "amino acid" refers to α -amino acids which are racemic, or of either the D- or L-configuration.

- As used herein, when any particular group, such as phenyl or pyridyl, is specified, this means that the group is unsubstituted or is substituted. Preferred substituents where not specified are halo, halo lower alkyl, and lower alkyl.
- 10

- As used herein, preparative or therapeutically useful amounts are amounts sufficient for use of the resulting products as therapeutics in which mgs to gms are administered. Typically, such amounts are at least 0.1 gms per synthesis, more preferably about gms to kilograms. Use of the LPCs provided herein results in improved yields and thereby permits development of preparative scale syntheses.
- 15

- As used herein, the abbreviations for any protective groups, amino acids and other compounds, are, unless indicated otherwise, in accord with their common usage, recognized abbreviations, or the IUPAC-IUB Commission on Biochemical Nomenclature (see, Biochem. 1972, 11, 1726).
- 20

A. Multifunctional Liquid Phase Carriers

- Liquid phase carriers for use in solution phase synthesis of biopolymers are provided. In particular, the LPCs provided herein are multifunctional LPCs that preferably possess three or more functional moieties for biopolymer synthesis. The biopolymers prepared on a given LPC may be of differing structure, but more typically are identical in structure.
- 25
- 30 The LPCs have the formula $Sp(X^1)_n$, wherein Sp is a polyvalent group that has more than two points of attachment, n corresponds to the

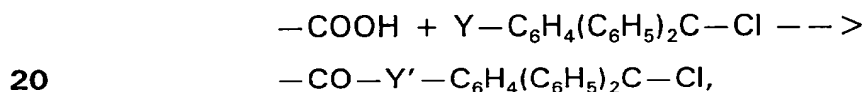
number of points of attachment in Sp and X^1 is a reactive group that is compatible from the point of view of biopolymer chemistry. Sp is preferably a symmetrical group such that all X^1 groups are equivalent. It will be appreciated by those of skill in the art that such symmetry allows

5 for uniformity in the synthesis of n biopolymers using these LPCs.

In preferred embodiments, multifunctional liquid phase carriers in which the groups X^1 can form a silyl ether, trityl ether or ester bond with the 3'- or 5'-hydroxyl groups of a nucleoside, nucleotide or oligonucleotide are used.

10 Examples of the reactive group X^1 which are customary in nucleotide chemistry are described in Liebigs Ann. Chem. **1978**, 839-853 and in Nucleic Acids Research, Symposium Series No. 7, **1980**, 39-59. Typical examples are, inter alia, the following:

1. Acid halides, in particular acid chlorides and acid bromides;
- 15 2. Carboxylic acid groups, which can react with 5'-OH groups, for example in the presence of condensing agents; they can also be converted into activated trityl chloride derivatives according to the following equation:



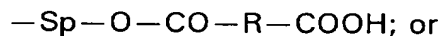
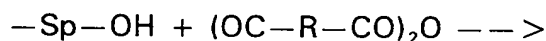
wherein $Y = OH, SH$ or NH_2 ; and $Y' = O, S$ or NH ;

3. Activated ester functions of the general formula $-COOR'$; and
4. OH, SH and NH_2 groups.

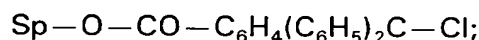
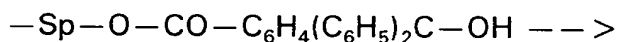
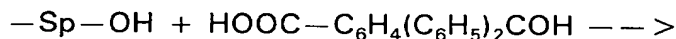
25 In another preferred embodiment, carrier molecules of the general formula $Sp(OH)_n$, and the corresponding thio and amino derivatives, are used directly in the provided methods if they are to react with reactive carboxyl groups. However, they are usually employed in the form of activated compounds, which can be obtained, for example, in the

30 following manner:

a) by reaction with dicarboxylic acid anhydrides, for example succinic or adipic anhydride, according to the equation



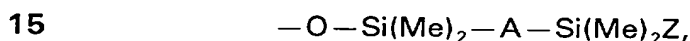
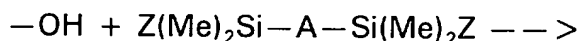
5 b) by reaction with trityl derivatives according to the equation



this reaction can be carried out in the presence of a condensing

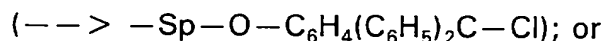
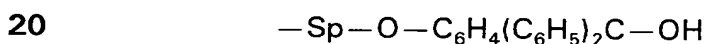
10 agent or after conversion of the acid function into an acid chloride function; or

c) by reaction with dichlorosiloxanes according to the general equation

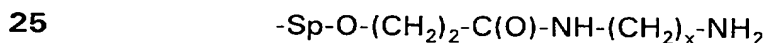


wherein A is alkylene, arylene, $-\text{O}-$ or a combination of these groups and Z is halogen; or

d) by reaction with hydroxyl-substituted trityl derivatives, to form trityl ether derivatives, for example of the general formula



e) by reaction with, for example, acrylonitrile, followed by alcoholysis of the nitrile and reaction of the resulting ester with an α,ω -diamine, to form amino derivatives, for example of the general formula



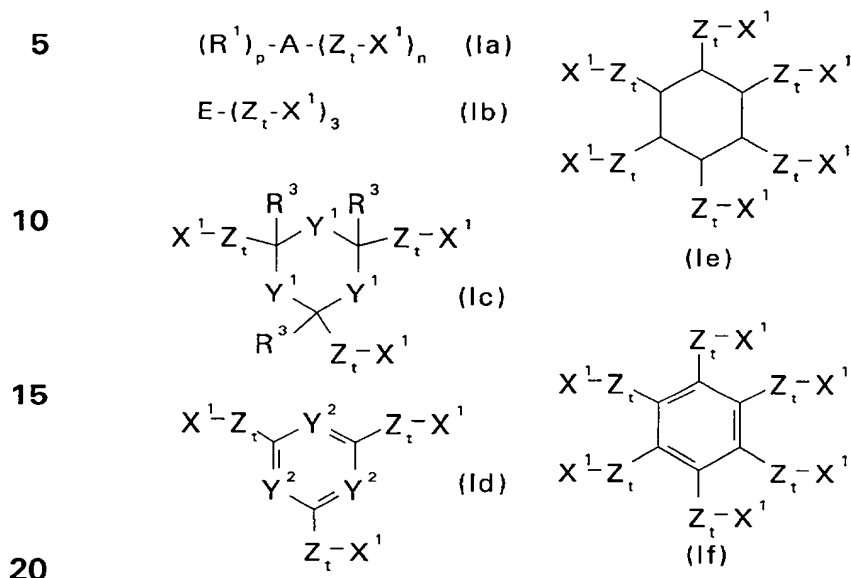
wherein x is 0-6. The amine is then further derivatized according to any of a) through d), above, prior to biopolymer synthesis.

If the corresponding thio or amino derivatives are used, the corresponding S or N compounds are formed.

30 In selecting the group Sp, it is particularly essential that the carrier molecule $\text{Sp}(\text{X}^1)_n$ and its derivatives obtained in the course of biopolymer

synthesis are soluble in the particular reaction medium used and can be chromatographed.

In one embodiment, the LPCs have formulae (I):



where A is carbon; E is nitrogen; R^1 and R^3 are each independently hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocyclylalkyl; p is 0 or 1; Z is a divalent hydrocarbon as described above, and is preferably any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12 units in which each unit is preferably selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X^1 is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably halide, OH, SH, NH_2 , COR^5 or $COOR^4$; n is 3 or 4; R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; R^5 is halide, heteroaryl or pseudohalide; Y^1 is CH_2 , NH, S or O; and Y^2 is selected from CH and N.

Any of the groups R^1 , R^3 , R^4 , R^5 , X^1 , Y^1 , Y^2 and Z may be unsubstituted or substituted with one or more substituents each

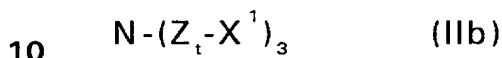
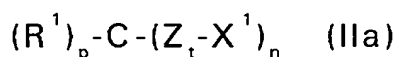
- independently selected from Q, which, as defined herein, is halogen, hydroxy, nitrile, nitro, formyl, mercapto, carboxy, alkyl, haloalkyl, polyhaloalkyl, aminoalkyl, diaminoalkyl, alkenyl containing 1 to 2 double bonds, alkynyl containing 1 to 2 triple bonds, cycloalkyl, cycloalkylalkyl,
- 5 aryl, heteroaryl, arylalkyl, heteroarylalkyl, alkylidene, arylalkylidene, alkylcarbonyl, arylcarbonyl, heteroarylcarbonyl, alkoxycarbonyl, alkoxycarbonylalkyl, aryloxycarbonyl, aryloxycarbonylalkyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, arylaminocarbonyl, diarylamino-
- 10 carbonyl, arylalkylaminocarbonyl, alkoxy, aryloxy, perfluoroalkoxy, alkenyloxy, alkynyloxy, arylalkoxy, amino, aminoalkyl, alkylaminoalkyl, dialkylaminoalkyl, arylaminoalkyl, diarylaminoalkyl, alkylamino, dialkylamino, arylamino, diarylamino, alkylarylamino, alkylcarbonylamino, alkoxycarbonylamino, arylcarbonylamino, aryloxycarbonylamino, azido, alkylthio, arylthio, perfluoroalkylthio, thiocyano, isothiocyano, alkyl-
- 15 sulfinyl, alkylsulfonyl, arylsulfinyl, arylsulfonyl, aminosulfonyl, alkylaminosulfonyl, dialkylaminosulfonyl, arylaminosulfonyl or diarylamino-sulfonyl.

- In embodiments where X^1 is halide, preferably chloride, Q is preferably selected from groups that are electron donating. Such groups
- 20 include alkyl, alkoxy, dialkylamino, diarylamino, alkylarylamino, alkylthio and arylthio, with alkoxy, preferably methoxy, being preferred.

- In certain embodiments, the LPCs are chosen as above, with the proviso that Z contains more than two methylene groups. In other
- embodiments, the LPCs are chosen with the proviso that Z is not 1,4-
- 25 arylene. In other preferred embodiments, the LPCs are chosen with the proviso that Z contains more than two methylene groups and is not 1,4-arylene. In other more preferred embodiments, the LPCs are chosen with the proviso that Z contains at least one, preferably at least two, more preferably at least three or more phenylene or alkylene units.

1. Acyclic Multifunctional LPCs

In one embodiment, the LPCs are acyclic, multifunctional compounds in which all of the functional groups are equivalent. These LPCs generally have a center of symmetry at a single atom. In particular,
 5 the LPCs have formulae (IIa) or (IIb):



where R^1 is hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocyclylalkyl; p and t are each independently 0 or 1; Z is any combination of phenylene and methylene units, selected from 0-12,
 15 preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; X^1 is any reactive group that is used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by
 20 reference), and is preferably OH, SH, NH_2 , COR^5 or $COOR^4$; n is 3 or 4; R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R^5 is halide, heteroaryl or pseudohalide.

In preferred embodiments, the LPCs are of formula (IIa) wherein p
 25 is 0 or 1, preferably 0, and n is 3 or 4, preferably 4. Thus in more preferred embodiments, the LPCs of formula (IIa) have the formula $C-(Z-X^1)_4$, where Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12, units selected from 1,4-phenylene and methylene,
 30 which units may be combined in any order; X^1 is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably OH, SH, NH_2 , COR^5 or $COOR^4$; R^4 is selected from

hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R^5 is halide, heteroaryl or pseudohalide.

In more preferred embodiments, the LPCs of formula (IIa) have the formula $C-(Z-X^1)_4$, where Z is C_{1-12} , preferably C_{2-12} , more preferably C_{3-12} , particularly C_{4-12} , most preferably C_{6-12} , C_{7-12} or C_{8-12} alkylene and X^1 is selected from OH, SH, NH_2 , COR^5 and $COOR^4$, where R^4 is hydrogen, alkyl, aryl or aralkyl and R^5 is halide, heteroaryl or pseudohalide.

In particularly preferred embodiments, the LPCs of formula (IIa) have the formula $C-(Z-X^1)_4$, where Z is C_{3-12} , particularly C_{4-12} , most preferably C_{6-12} , C_{7-12} or C_{8-12} alkylene and X^1 is selected from OH, SH, NH_2 , COR^5 and $COOR^4$, where R^4 is hydrogen, alkyl, aryl or aralkyl and R^5 is halide, heteroaryl or pseudohalide. In more preferred embodiments, X^1 is OH, SH or NH_2 , preferably OH.

In other particularly preferred embodiments, the LPCs of formula (IIa) have the formula $C-(Z-X^1)_4$, where Z is methylene or ethylene, preferably methylene, and X^1 is selected from OH, SH, NH_2 , COR^5 and $COOR^4$, where R^4 is hydrogen, alkyl, aryl or aralkyl and R^5 is halide, heteroaryl or pseudohalide. In more preferred embodiments, X^1 is OH, SH or NH_2 , preferably OH. The presently most preferred LPC of formula (IIa) is pentaerythritol.

In other preferred embodiments, the LPCs are of formula (IIb) where Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; X^1 is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably OH, SH, NH_2 , COR^5 or $COOR^4$; R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R^5 is halide, heteroaryl or pseudohalide.

In more preferred embodiments, the LPCs have formula (IIb), where Z is C₁₋₁₂, preferably C₂₋₁₂, more preferably C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide.

In particularly preferred embodiments, the LPCs have formula (IIb), where Z is C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is OH, SH or NH₂, preferably NH₂.

In other particularly preferred embodiments, the LPCs have formula (IIb), where Z is methylene or ethylene, preferably ethylene, and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is OH, SH or NH₂, preferably NH₂. The presently most preferred LPC of formula (IIb) is tri(2-aminoethyl)amine.

In certain embodiments, the LPCs of formulae (IIa) and (IIb) are chosen as above, with the proviso that Z contains more than two, preferably more than three, more preferably more than four methylene groups. In other embodiments, the LPCs are chosen with the proviso that Z is not 1,4-arylene. In other preferred embodiments, the LPCs are chosen with the proviso that Z contains more than two methylene groups and is not 1,4-arylene. In other more preferred embodiments, the LPCs are chosen with the proviso that Z contains at least one, preferably at least two, more preferably at least three or more phenylene or alkylene units.

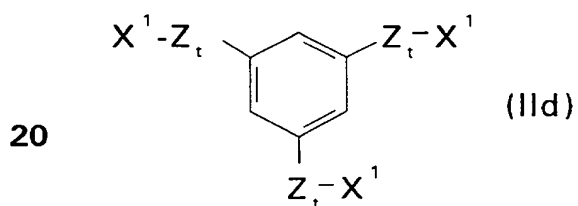
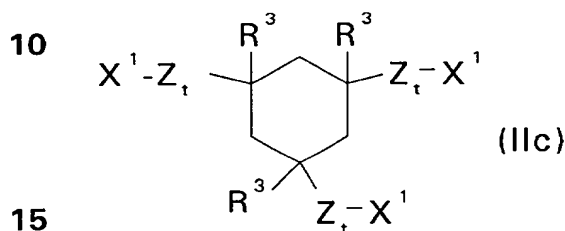
2. Cyclic Multifunctional LPCs

In another embodiment, the LPCs are cyclic, multifunctional compounds in which all of the functional groups are equivalent. These LPCs generally have a center of symmetry at the center of a core

5 cycloalkyl, heterocyclyl, aryl or heteroaryl ring.

a. Cyclic Multifunctional LPCs of Formulae (IIc) or (IId)

In more preferred embodiments, the LPCs have formulae (IIc) or (IId):



where Z is any combination alkylene and phenylene units, of 0-12 units,
 25 preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12 units, preferably selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which
 30 is incorporated herein by reference), and is preferably halide, OH, SH, NH₂, COR⁵ or COOR⁴; n is 3 or 4; R⁴ is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R⁵ is halide, heteroaryl or pseudohalide.

In other preferred embodiments, the LPCs are of formula (IIc) where
 35 Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-

12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is

- 5 preferably halide, OH, SH, NH₂, COR⁵ or COOR⁴; R⁴ is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclalkyl; and R⁵ is halide, heteroaryl or pseudohalide.

- In more preferred embodiments, the LPCs have formula (IIc), where Z is C₁₋₁₂, preferably C₂₋₁₂, more preferably C₃₋₁₂, particularly C₄₋₁₂, most
10 preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene; t is 1; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide.

- In particularly preferred embodiments, the LPCs have formula (IIc), where Z is C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂
15 alkylene; t is 1; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is COR⁵ or COOR⁴, preferably COOR⁴.

- In other particularly preferred embodiments, the LPCs have formula
20 (IIc), where Z contains 0, 1 or 2 methylene units, preferably 0; t is preferably 0; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is COR⁵ or COOR⁴, preferably COOR⁴.

- 25 In other preferred embodiments, the LPCs are of formula (IId) where Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can
30 be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is

preferably halide, OH, SH, NH₂, COR⁵ or COOR⁴; R⁴ is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R⁵ is halide, heteroaryl or pseudohalide.

In more preferred embodiments, the LPCs have formula (IId), where

5 Z is C₁₋₁₂, preferably C₂₋₁₂, more preferably C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene; t is 1; and X¹ is selected from halide, preferably chloride, OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide.

10 In particularly preferred embodiments, the LPCs have formula (IId), where Z is C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene; t is 1; and X¹ is selected from halide, preferably chloride, OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred
15 embodiments, X¹ is halide or COR⁵ or COOR⁴, preferably chloride or COOR⁴.

In other particularly preferred embodiments, the LPCs have formula (IId), where Z contains 0, 1 or 2 methylene groups, preferably 0 (*i.e.*, Z is absent); t is preferably 0; and X¹ is selected from halide, preferably

20 chloride, OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl, aryl or pseudohalide. In more preferred embodiments, X¹ is halide, COR⁵ or COOR⁴, preferably chloride or COOR⁴. The presently most preferred LPCs of formula (IId) are 1,3,5-benzenetricarboxylic acid and 1,3,5-tri(chloromethyl)benzene.

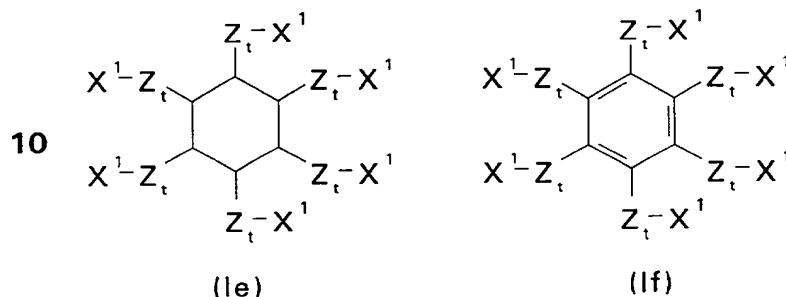
25 In certain embodiments, the LPCs of formulae (IIc) and (IId) are chosen as above, with the proviso that Z contains more than two methylene groups. In other embodiments, the LPCs are chosen with the proviso that Z is not 1,4-arylene. In other preferred embodiments, the LPCs are chosen with the proviso that Z contains more than two
30 methylene groups and is not 1,4-arylene. In other more preferred embodiments, the LPCs are chosen with the proviso that Z contains at

least one, preferably at least two, more preferably three or more phenylene or methylene units.

b. Cyclic Multifunctional LPCs of Formulae (Ie) and (If)

In other more preferred embodiments, the LPCs have formulae (Ie)

5 or (If):



15

where Z is any combination of alkylene or arylene units as described above, of 0-12 units, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12 units

20 preferably selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably OH, SH, NH₂, COR⁵ or COOR⁴; n is 3 or 4; R⁴ is selected from

25 hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R⁵ is halide, heteroaryl or pseudohalide.

In other preferred embodiments, the LPCs are of formula (Ie) where Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-

30 12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably OH, SH, NH₂, COR⁵ or COOR⁴; R⁴ is selected from hydrogen,

alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R⁵ is halide, heteroaryl or pseudohalide.

In more preferred embodiments, the LPCs have formula (Ie), where Z is C₁₋₁₂, preferably C₂₋₁₂, more preferably C₃₋₁₂, particularly C₄₋₁₂, most
 5 preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene; t is 1; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide.

In particularly preferred embodiments, the LPCs have formula (Ie), where Z is C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂
 10 alkylene; t is 1; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is COR⁵ or COOR⁴, preferably COOR⁴.

In other particularly preferred embodiments, the LPCs have formula
 15 (Ie), where Z contains 0, 1 or 2 methylene units, preferably 0; t is preferably 0; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is COR⁵ or COOR⁴, preferably COOR⁴, most preferably COOH.

In other preferred embodiments, the LPCs are of formula (If) where
 20 Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can
 25 be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably OH, SH, NH₂, COR⁵ or COOR⁴; R⁴ is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R⁵ is halide, heteroaryl or pseudohalide.

In more preferred embodiments, the LPCs have formula (If), where
 30 Z is C₁₋₁₂, preferably C₂₋₁₂, more preferably C₃₋₁₂, particularly C₄₋₁₂, most

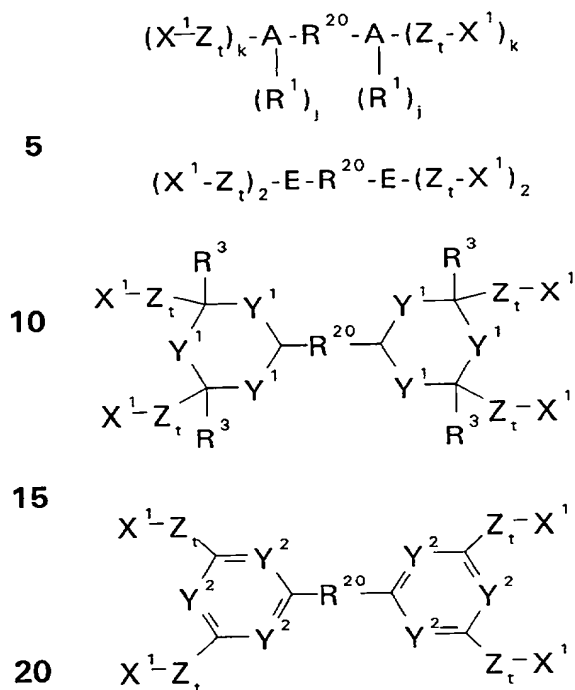
preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene; t is 1; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide.

- In particularly preferred embodiments, the LPCs have formula (If),
- 5 where Z is C₃₋₁₂, particularly C₄₋₁₂, most preferably C₆₋₁₂, C₇₋₁₂ or C₈₋₁₂ alkylene; t is 1; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl or pseudohalide. In more preferred embodiments, X¹ is COR⁵ or COOR⁴, preferably COOR⁴.
- 10 In other particularly preferred embodiments, the LPCs have formula (If), where Z contains 0, 1 or 2 methylene groups, preferably 0 (*i.e.*, Z is absent); t is preferably 0; and X¹ is selected from OH, SH, NH₂, COR⁵ and COOR⁴, where R⁴ is hydrogen, alkyl, aryl or aralkyl and R⁵ is halide, heteroaryl, aryl or pseudohalide. In more preferred embodiments, X¹ is
- 15 COR⁵ or COOR⁴, preferably COOR⁴, most preferably COOH.

- In certain embodiments, the LPCs of formulae (Ie) and (If) are chosen as above, with the proviso that Z contains more than two methylene groups. In other embodiments, the LPCs are chosen with the proviso that Z is not 1,4-arylene. In other preferred embodiments, the
- 20 LPCs are chosen with the proviso that Z contains more than two methylene groups and is not 1,4-arylene. In other more preferred embodiments, the LPCs are chosen with the proviso that Z contains at least one, preferably at least two, more preferably three or more phenylene or methylene units.

25 3. Dimeric LPCs

- In preferred embodiments, the LPCs are composed of two of the groups of formula (I). The LPCs of this embodiment are symmetrical compounds such that all of the X¹ groups are equivalent. In more
- 30 preferred embodiments, the LPCs have formulae:



- where A is carbon; E is nitrogen; R¹ and R³ are each independently hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocyclylalkyl; p is 0 or 1; Z as defined above is any combination of 0-12 units, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12 units selected from 1,4-phenylene and methylene, which units may be combined in any order; R²⁰ is alkylene; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably OH, SH, NH₂, COR⁵ or COOR⁴; n is 3 or 4; R⁴ is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; R⁵ is halide, heteroaryl or pseudohalide; Y¹ is CH₂, NH, S or O; and Y² is selected from CH and N.

In more preferred embodiments, Y^1 is CH_2 and Y^2 is CH .

Any of the groups R^1 , R^3 , R^4 , R^5 , X^1 , Y^1 , Y^2 and Z may be unsubstituted or substituted with one or more substituents each independently selected from Q .

- 5 In other preferred embodiments, the LPCs are of the above formulae where Z is any combination of 0-12, preferably 1-12, more preferably 2-12, particularly 3-12, more particularly 4-12, most preferably 6-12, 7-12 or 8-12, units selected from 1,4-phenylene and methylene, which units may be combined in any order; t is 0 or 1; R^{20} is C_{1-12} , preferably C_{1-6} , alkylene; X^1 is any reactive group which can be used in biopolymer synthesis (see, e.g., U.S. Patent No. 5,198,540, the disclosure of which is incorporated herein by reference), and is preferably halide, OH , SH , NH_2 , COR^5 or $COOR^4$; R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R^5 is halide, heteroaryl or pseudohalide.
- 10
- 15

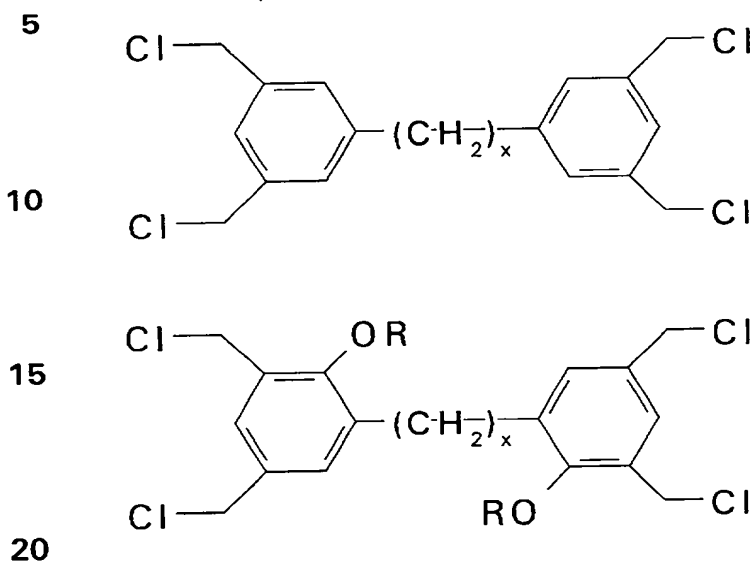
- In more preferred embodiments, the LPCs have the above formulae, where Z is C_{1-12} , preferably C_{2-12} , more preferably C_{3-12} , particularly C_{4-12} , most preferably C_{6-12} , C_{7-12} or C_{8-12} alkylene; t is 1; R^{20} is lower alkylene; and X^1 is selected from halide, OH , SH , NH_2 , COR^5 and $COOR^4$, where R^4 is hydrogen, alkyl, aryl or aralkyl and R^5 is halide, heteroaryl or pseudohalide.
- 20

- In particularly preferred embodiments, the LPCs have the above formulae, where Z is C_{3-12} , particularly C_{4-12} , most preferably C_{6-12} , C_{7-12} or C_{8-12} alkylene; t is 1; R^{20} is lower alkylene; and X^1 is selected from OH , SH , NH_2 , COR^5 and $COOR^4$, where R^4 is hydrogen, alkyl, aryl or aralkyl and R^5 is halide, heteroaryl or pseudohalide. In more preferred embodiments, X^1 is COR^5 or $COOR^4$, preferably $COOR^4$.
- 25

- In other particularly preferred embodiments, the LPCs have formula (IIId), where Z contains 0, 1 or 2 methylene groups, preferably 0 (i.e., Z is absent) or 1; t is 0 or 1; R^{20} is lower alkylene; and X^1 is selected from halide, preferably chloride, OH , SH , NH_2 , COR^5 and $COOR^4$, where R^4 is
- 30

hydrogen, alkyl, aryl or aralkyl and R^5 is halide, heteroaryl, aryl or pseudohalide. In more preferred embodiments, X^1 is halide, COR^5 or $COOR^4$, preferably chloride or $COOR^4$, most preferably chloride or $COOH$.

Most preferred compounds of the above formulae include:



where x is 0-6 and R is lower alkyl.

4. LPC Derivatives

In certain embodiments, it is desirable to modify the LPC prior to biopolymer synthesis. Such modification produces an LPC that has desirable physical characteristics, such as solubility or compatibility with chromatography, for biopolymer synthesis. The type of modification to be performed will depend on the nature of the biopolymer to be prepared.

a. LPCs for Oligonucleotide Synthesis

LPCs for use in oligonucleotide synthesis will preferably possess a carboxylic acid group or a derivative thereof at the terminus of X^1 . This type of functionality may be used to form an ester linkage to, for example, the 3'-OH group of a first nucleotide or nucleoside monomer.

If X^1 is OH or SH, the modification involves reaction with, for example, acrylonitrile, followed by alcoholysis of the nitrile and reaction of the resulting ester with an α,ω -diamine, to form an amino derivative, for example, of the general formula $Sp(O,S-(CH_2)_2-C(O)-NH-(CH_2)_x-NH_2)_n$

where x is 0-6, preferably 1-4, more preferably 2. The presently most preferred LPC of this formula is tetrakis(8-amino-6-aza-2-oxa-5-oxooctyl)methane. The amine may then be further derivatized by reaction with, for example, an α,ω -alkanedioic acid anhydride to provide a compound of formula $\text{Sp}(\text{O},\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, where x is 0-6, preferably 1-4, more preferably 2. The presently most preferred LPC of this formula is tetrakis(11-carboxy-6,9-diaza-5,10-dioxo-2-oxaundecyl)methane.

In embodiments where X^1 is NH_2 , the LPC may be derivatized by reaction with, for example, an α,ω -alkanedioic acid anhydride to provide a compound of formula $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, where x is 0-6, preferably 1-4, more preferably 2. The presently most preferred LPC of this formula is tris(3-aza-6-carboxy-4-oxohexyl)amine.

In embodiments where X^1 is COOR^4 or COR^5 , the LPC may be derivatized by reaction with, for example, an α,ω -alkylenediamine, to form an amino derivative, for example, of the general formula $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}_2)_n$ where x is 0-6, preferably 1-4, more preferably 2. The presently most preferred LPC of this formula is 1,3,5-benzenetricarboxylic acid tris-N-(2-aminoethyl)amide. The amine may then be further derivatized by reaction with, for example, an α,ω -alkanedioic acid anhydride to provide a compound of formula $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, where x is 0-6, preferably 1-4, more preferably 2. The presently most preferred LPC of this formula is 1,3,5-benzenetricarboxylic acid tris-N-(3-aza-6-carboxy-4-oxohexyl)amide.

In other more preferred embodiments, the carboxyl derivatives of formulae $\text{Sp}(\text{O},\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$ and $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$ may be further derivatized by reaction with a first monomer unit for synthesis of a biopolymer. As exemplified herein, a first monomer unit in the synthesis of oligonucleotides may be 5'-O-dimethoxytrityl-2'-deoxythymidine or 2'-deoxythymidine. Preferred LPCs of this

- embodiment include tetrakis{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-dT)₄-PE-LPC), 1,3,5-tris{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenyl-methyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC), tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC), tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine ((DMT-dT)₃-Amine-LPC) and tris[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

b. LPCs for Peptide Synthesis

- LPCs for use in peptide synthesis will preferably possess a haloalkyl, hydroxyl, thio or carboxyl group at the terminus of X¹. The hydroxyl or thio groups may be used to form ester or thioester linkages to the carboxyl group of a first amino acid monomer. Preferred haloalkyl groups are benzylic halides that can form benzylic esters with the carboxyl terminus of a first monomer. The carboxyl group of X¹ may be used to form an amide linkage to the amino group of a first amino acid monomer. LPCs possessing carboxyl groups at the terminus of X¹ are described above.

- In embodiments where X¹ is OH or SH, the modification involves reaction with, for example, acrylonitrile, followed by alcoholysis of the nitrile and reaction of the resulting ester with an α,ω -diol, dithiol or oxythiol, to form a hydroxy or thio derivative, for example, of the general formula $\text{Sp}(\text{O},\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{O},\text{S}-(\text{CH}_2)_x-\text{OH},\text{SH})_n$ where x is 0-6, preferably 1-4, more preferably 2.

- In embodiments where X¹ is NH₂, the LPC may be derivatized by reaction with, for example, an α,ω -alkanedioic acid anhydride to provide a compound of formula $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, where x is 0-6, preferably 1-4, more preferably 2. Reaction of the resulting acid with an

α,ω -diol, dithiol or oxy-thiol, forms a hydroxy or thio derivative, for example, of the general formula $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{CO}-\text{O},\text{S}-(\text{CH}_2)_x-\text{OH},\text{SH})_n$, where x is 0-6, preferably 1-4, more preferably 2.

- In embodiments where X^1 is COOR^4 or COR^5 , the LPC may be
- 5 derivatized by reaction with, for example, an α,ω -diol, dithiol or oxy-thiol, to form a hydroxy or thio derivative, for example, of the general formula $\text{Sp}(\text{C}(\text{O})-\text{O},\text{S}-(\text{CH}_2)_x-\text{OH},\text{SH})_n$, where x is 0-6, preferably 1-4, more preferably 2.

- In other more preferred embodiments, the hydroxy or thio
- 10 derivatives of formulae $\text{Sp}(\text{O},\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{O},\text{S}-(\text{CH}_2)_x-\text{OH},\text{SH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{CO}-\text{O},\text{S}-(\text{CH}_2)_x-\text{OH},\text{SH})_n$ or $\text{Sp}(\text{C}(\text{O})-\text{O},\text{S}-(\text{CH}_2)_x-\text{OH},\text{SH})_n$ may be further derivatized by reaction with a first monomer unit for synthesis of a biopolymer.

c. LPCs for Peptide Nucleic Acid (PNA) Synthesis

- 15 LPCs for use in PNA synthesis will preferably possess a haloalkyl, hydroxyl, thio or carboxyl group at the terminus of X^1 . The hydroxyl or thio groups may be used to form ester or thioester linkages to the carboxyl group of a first PNA monomer. Preferred haloalkyl groups are benzylic halides that can form benzylic esters with the carboxyl terminus
- 20 of a first monomer. The carboxyl group of X^1 may be used to form an amide linkage to the amino group of a first PNA monomer. LPCs possessing these groups at the terminus of X^1 are described above.

d. LPCs for Oligosaccharide Synthesis

- LPCs for use in oligosaccharide synthesis will preferably possess a
- 25 hydroxyl, thio or carboxyl group at the terminus of X^1 . The hydroxyl or thio groups may be used to form glycosyl linkages to the a first saccharide monomer. The carboxyl group may be used to form an ester linkage to a hydroxyl (e.g., the 5- or 6-hydroxyl) group of a first saccharide monomer. LPCs possessing these groups at the terminus of
- 30 X^1 are described above.

e. LPCs Coupled to Linkers

- Alternatively, or in conjunction with the modifications described above, the LPC may be coupled with any linker known to those of skill in the art for use in biopolymer synthesis. Such linkers have traditionally
- 5 been used in solid phase synthesis of biopolymers. The linkers include photocleavable, traceless, safety-catch or other linkers well known to those of skill in the art. See, e.g., Leznoff *et al.* Can. J. Chem. **1972**, 50, 2892-2893; Frechet *et al.* Can. J. Chem. **1976**, 54, 926-934; Chan *et al.* J. Chem. Soc., Chem. Commun. **1995**, 1475-1476; Krchnak *et al.* Mol.
 - 10 Divers. **1995**, 1, 149-164; Burgess *et al.* J. Org. Chem. **1997**, 62, 5165-5168; Plunkett *et al.* J. Org. Chem. **1995**, 60, 6006-6007; Han *et al.* Tetrahedron Lett. **1996**, 37, 2703-2706; Lorbach *et al.* J. Org. Chem. **1996**, 61, 8716-8717; Newlander *et al.* J. Org. Chem. **1997**, 62, 6726-6732; Worster *et al.* Angew. Chem. Int. Ed. Eng. **1979**, 18, 221-222;
 - 15 Allin *et al.* Tetrahedron Lett. **1996**, 37, 8023-8026; Lloyd-Williams *et al.* Tetrahedron **1993**, 49, 11065-11133; Brown *et al.* Mol. Divers. **1995**, 1, 4-12; Salmon *et al.* Proc. Natl. Acad. Sci. U.S.A. **1993**, 90, 11708-11712; Cardno *et al.* Tetrahedron Lett. **1996**, 37, 135-138; Kaldor *et al.* Tetrahedron Lett. **1996**, 37, 7193-7196; DeGrado *et al.* J. Org. Chem.
 - 20 **1980**, 45, 1295-1300; Bhargava *et al.* J. Am. Chem. Soc. **1983**, 105, 3247-3251; Van Maarseveen *et al.* Tetrahedron Lett. **1996**, 37, 8249-8252; Entwistle *et al.* Tetrahedron Lett. **1979**, 555-558; Maeji *et al.* J. Immunol. Meth. **1990**, 134, 23-33; Bray *et al.* J. Org. Chem. **1991**, 56, 6659-6666; Backes *et al.* J. Am. Chem. Soc. **1996**, 118, 3055-3056;
 - 25 Gayo *et al.* Tetrahedron Lett. **1997**, 38, 211-214; Morphy *et al.* Tetrahedron Lett. **1996**, 37, 3209-3212, Patek *et al.* Tetrahedron Lett. **1991**, 32, 3891-3894; Routledge *et al.* Tetrahedron Lett. **1997**, 38, 1227-1230; Fattom *et al.* (1992) Infection & Immun. **60**:584-589; Goldmacher *et al.* (1992) Bioconj. Chem. **3**:104-107; Hazum *et al.* (1981)
 - 30 in Pept., Proc. Eur. Pept. Symp., 16th, Brunfeldt, K (Ed), pp. 105-110; Yen *et al.* (1989) Makromol. Chem **190**:69-82; Senter *et al.* (1985)

Photochem. Photobiol 42:231-237 and U.S. Patent Nos. 5,547,835, 5,635,598, 5,492,821, 5,652,358, 5,552,471, 5,736,626, 5,639,633, 5,736,625, 5,679,773, 5,648,462, 5,616,698 and 5,552,535. The disclosure of these references is herein incorporated by reference.

5 B. Preparation of Liquid Phase Carriers

The liquid phase carriers may be prepared by the methods described in detail herein, by minor modification of the methods or by other methods known to those of skill in the art. Certain of the LPCs described herein are commercially available from, for example, Aldrich

10 Chemical Co., Milwaukee, WI.

1. Preparation of LPCs of Formula (Ia)

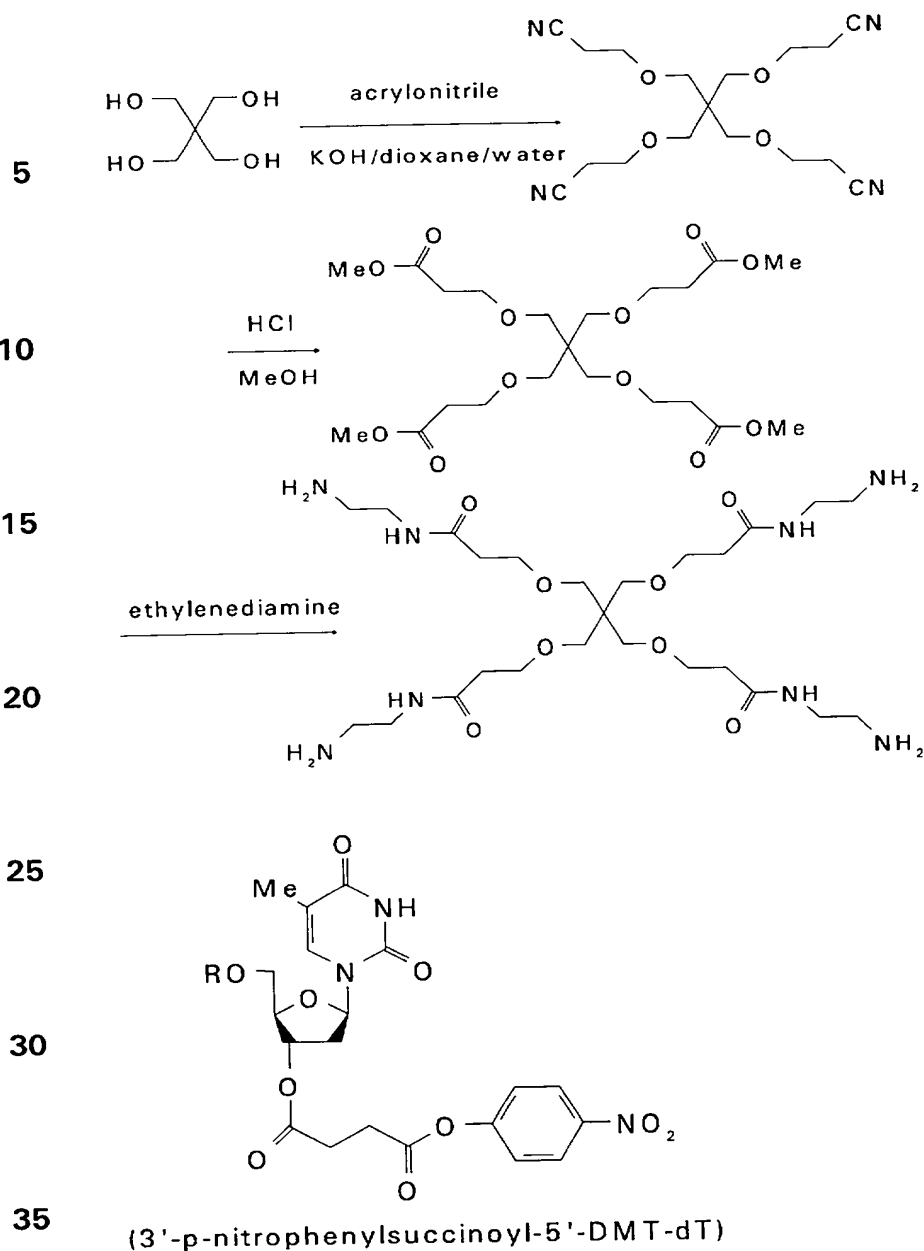
Preparation of LPCs of formula (Ia) is exemplified by the syntheses of tetrakis{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-
15 dT)₄-PE-LPC) and tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), shown below (see also, Examples 1 and 3). Other LPCs of formula (Ia) may be prepared by minor modification of this method.

20

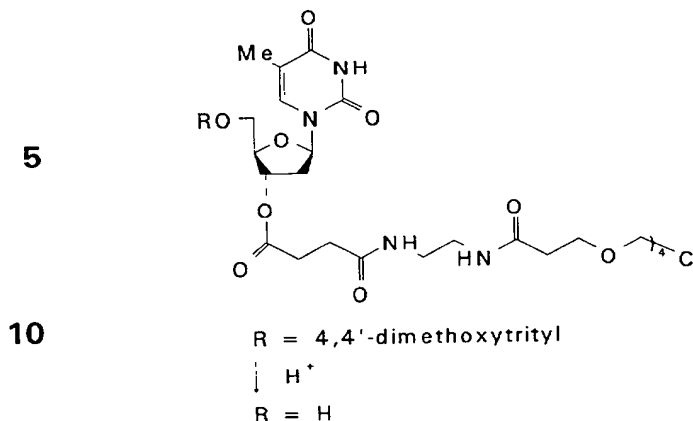
25

30

35



pyridine
R = 4,4'-dimethoxytrityl



- 15 Treatment of pentaerythritol with acrylonitrile in the presence of KOH gave tetrakis-[(cyanoethoxy)methyl]methane. When refluxed with HCl saturated methanol under exclusion of water the nitrile reacts to form tetrakis-[(methoxy-carbonyl)ethoxy)methyl]methane. See, G.R. Newkome *et al.* *Aldrichim. Acta* **1992**, 25, 31-38; G.R. Newkome *et al.*
- 20 *Macromolecules* **1991**, 24, 1443-1444 and G.R. Newkome *et al.* *Tetrahedron: Asymmetry*, **1991**, 2, 957-960. Aminolysis with ethylendi-amine leads to the tetravalent amine, tetrakis-(8-amino-6-aza-2-oxa-5-oxooctyl)-methane. See, A.D. Meltzer *et al.* *Macromolecules* **1992**, 25, 4541-4548. The four primary amino functions were then acylated by the
- 25 reactive p-nitrophenylester of 5'-O-dimethoxytrityl-deoxythymidine-3'-O-succinate in pyridine, leading to the highly symmetrical compound tetrakis-{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-dT)₄-PE-LPC). Repeated low pressure liquid chromatography on silica gel
- 30 and gel permeation chromatography (GPC) on Sephadex LH20 afforded pure material in a yield of 45%.

The next step, the dedimethoxytritylation of the LPC is the first necessary reaction for starting the synthesis of oligonucleotides and is also repeated at the beginning of every following cycle. An important

35 aspect of this procedure is that the deprotection of the terminal 5'-hydroxyl group be rapid and complete. Long reaction times under acidic

conditions will favor another known side reaction, the depurination of guanosine and adenosine units in the growing oligonucleotide chain.

In one method, cationic ion exchanger resins in their protonated form (Fractogel® EMD-SO₃⁻ 650 (M) in H⁺-form) may be used for combined
 5 dedimethoxytritylation and purification by stepwise variation of the polarity of the solvent system (solid phase extraction). Nonpolar solvents (dichloromethane/ethanol mixtures) enable the fast separation of the dimethoxytrityl group with high flow rates. The 5'-O-deprotected LPC is eluted next with polar solvent systems such as ethanol or THF/ethanol
 10 mixtures. The yield of dT₄-PE-LPC was 93%.

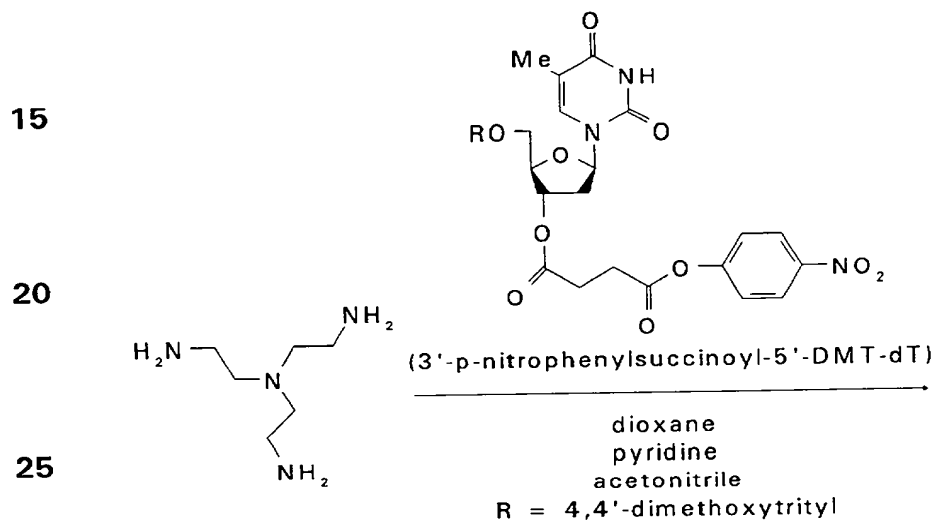
In another method, dedimethoxytritylation with TFA (2%) in 1,2-dichloroethane/nitromethane/methanol 80:19:1 (v/v) was used for rapid cleavage of the 5'-O-protecting group, completed in less than 5 minutes. After neutralization (with, for example, triethylamine), purification may be
 15 performed with different types of chromatography media:
 a) by solid phase extraction using LiChroprep® DIOL, or
 b) by GPC using Sephadex LH20 and additional extraction of remaining dimethoxytritylcarbinol with diethyl ether, or
 c) neutralization with n-trioctylamine, purification by reversed phase
 20 chromatography using NUCLEOPREP® 300-30 C18 (eluant: acetonitrile/THF/water 56:14:30 v/v), or
 d) by reversed phase chromatography using NUCLEOPREP® 300-30 C18 in combination with gel permeation chromatography using Sephadex LH20 eluant: THF/water 70:30 v/v.

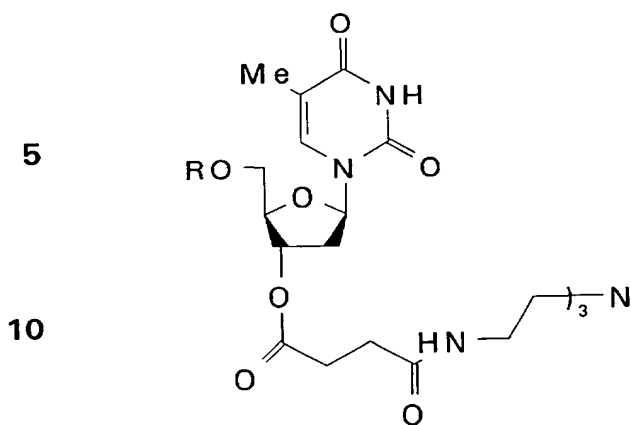
25 The use of water in these methods makes intensive drying of nucleotidic products necessary before the following coupling reaction with phosphoramidites is started. In general water is not necessary eluent for chromatography using only Sephadex LH20. Only a few very polar solvents are able to fulfill the demands for good solubility of all
 30 components and for other properties in this chromatography (sufficient swelling of Sephadex LH20, low viscosity, low boiling points for easy

removal after GPC). Pyridine is preferred for this purpose, also because this solvent proved compatible with the condensation reaction with phosphoramidites for chain elongation, in contrast to water, alcohols, organic acids or strong bases.

5 2. Preparation of LPCs of Formula (Ib)

Preparation of LPCs of formula (Ib) is exemplified by the synthesis of tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine ((DMT-dT)₃-Amine-LPC) and tris-[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC), shown below (see also, Examples 5 and 6). Other LPCs of formula (Ib) may be prepared by minor modification of this method.





15

R = 4,4'-dimethoxytrityl
 \downarrow H⁺
 R = H

20

This LPC may be prepared by minor modification of the method described above for synthesis of LPCs of formula (Ia). Synthesis starting with tris(2-aminoethyl)amine provided (DMT-dT)₃-Amine-LPC in good yield (74%). Dedimethoxytritylation under conditions described for LPCs of formula (Ia) gave dT₃-Amine-LPC.

25

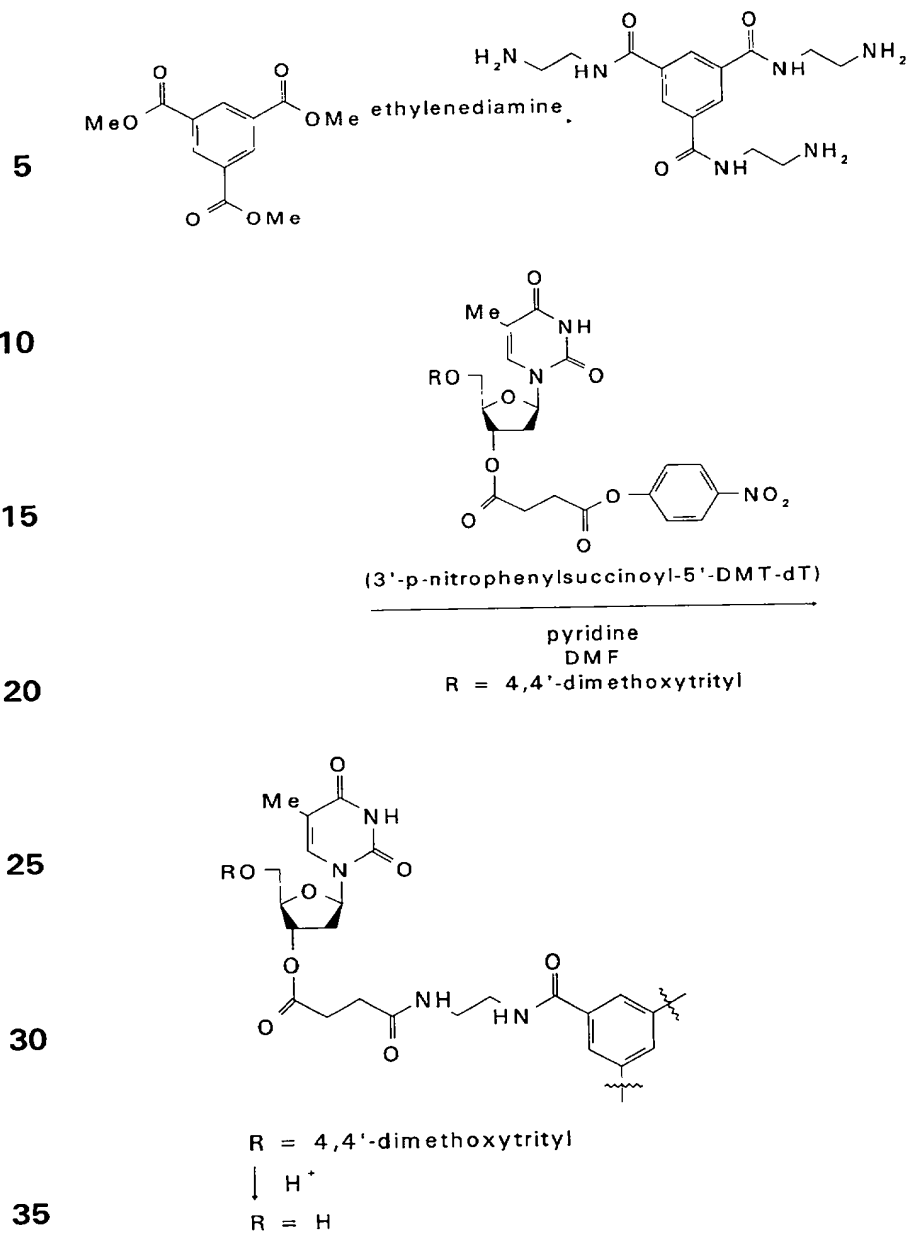
3. Preparation of LPCs of Formulae (Ic) and (Id)

Preparation of LPCs of formulae (Ic) and (Id) are exemplified by the synthesis of 1,3,5-tris-{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC) and 1,3,5-tris-[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC), shown below (see also, Examples 2 and 4). Other LPCs of formulae (Ic) and (Id) may be prepared by minor modification of this method.

30

35

40



Aminolysis of commercially available 1,3,5-benzene tricarboxylic acid trimethyl ester using ethylenediamine gave the triamide shown. This triamide, 1,3,5-benzene tricarboxylic acid tris-N-(2-aminoethyl)amide, was then treated with the p-nitrophenylester of 5'-O-dimethoxytrityl-deoxythymidine-3'-O-succinate in pyridine/DMF to give an aryl type liquid phase carrier, abbreviated as (DMT-dT)₃-Aryl-LPC, in yields of up to 90%.

Silica gel column chromatography and GPC on Sephadex LH20 were used for purification, ^1H - and ^{13}C -NMR for identification.

C. Methods of Solution Phase Biopolymer Synthesis

Methods of synthesis of biopolymers using the LPCs described herein are also provided. Relatively large amounts of biopolymers, including, but not limited to, oligonucleotides, peptides, PNAs and oligosaccharides, may be produced by the methods. It is possible to prepare not only naturally occurring biopolymers, but also analogs of the biopolymers. Such analogs include, for example, analogs of oligonucleo-
 10 tides such as oligo-triesters or oligophosphonates. Use of the LPCs provided herein in the methods provided afford biopolymers of high purity and in significantly higher yields than methods known in the art.

In one embodiment, the methods involve the steps of (a) reacting an LPC of formula $\text{Sp}(\text{X}^1)_n$, provided herein, with a first monomer N^1 ; (b)
 15 separating and purifying the product of step (a) to afford a compound of formula $\text{Sp}(\text{X}^1\text{-N}^1)_n$; (c) reacting the product of step (b) with a second monomer N^2 , a dimer $\text{N}^2\text{-N}^3$ or trimer $\text{N}^2\text{-N}^3\text{-N}^4$; (d) repeating steps (b) and (c) using the desired monomers, dimers or trimers to produce the desired LPC-bound biopolymer of formula $\text{Sp}(\text{X}^1\text{-N}^1\text{-N}^2\text{-...-N}^m)_n$, where m is 3 to
 20 100, preferably 3-50, more preferably 3-30, most preferably 3-15 or 3-10; and (e) cleaving the biopolymer from the LPC.

In embodiments where the biopolymer is an oligonucleotide, peptide or a PNA, m is preferably 3-30, more preferably 3-15. In
 25 3-15, more preferably 3-10. In embodiments where the biopolymer is an oligosaccharide, m is preferably

The separation and purification steps may be accomplished by any method known to those of skill in the art. Preferred methods include gel permeation chromatography (GPC) and chromatography on reverse phase (RP) silica gel. In more preferred embodiments, combinations of GPC and
 30 RP-silica gel chromatography are used.

Cleavage of the biopolymer from the LPC is achieved through standard means and depends on the nature of the linkage between the LPC and the biopolymer. For example, ester linkages may be cleaved with a base, such as hydroxide, alkoxides and amines. In certain
 5 embodiments, ammonium hydroxide is preferred. In other embodiments, the linkages may be silyl or trityl ethers, or glycosyl bonds. In these embodiments, treatment with acid, such as acetic acid, will cleave the biopolymer from the LPC. In embodiments where a linker is used to couple the LPC to the biopolymer, the cleavage conditions will depend on
 10 the nature of the linker. For example, light of a specific wavelength can be used to release a biopolymer from a photocleavable linker.

1. Monomers, Dimers and Trimers

Monomers (N), dimers and trimers used in preparing the biopolymer will depend on the biopolymer desired. For example, monomers which
 15 are used in the methods include nucleotides, nucleosides, natural and unnatural amino acids, PNA monomers and monosaccharides. Dimers and trimers are composed of the monomers. The monomers are used or obtained in the course of synthesis in protected form; typical examples of such protective groups are well known to those of skill in the art (see,
 20 e.g., Tetrahedron **1981**, 363-369; Liebigs Ann. Chem. **1978**, 839-853; Nucleic Acids Research, Symposium Series No. 7 **1980**, 39-59; and T. W. Greene Protective groups in Organic Synthesis (John Wiley & Sons, Inc., 1981)).

2. Biopolymeric Synthetic Protocols

25 In the methods, synthesis of the biopolymeric chain (i.e., steps c and d) may be carried out using any protocols known to those of skill in the art. For example, in the synthesis of oligonucleotides, the phosphate triester, phosphite triester, H-phosphonate or phosphoramidite protocols may be used. See, e.g., N. D. Sinha et al. Tetrahedron Lett. **1983**, 24,
 30 5843-5846; H. Köster et al. Tetrahedron **1984**, 40, 103-112; N.D. Sinha et al. Nucleic Acids Res. **1984**, 12, 4539-4557; S.L. Beaucage et al.

- Tetrahedron **1992**, 48, 2223-2311; E.Sonveaux Bioorg. Chem. **1986**, 14, 274-325; P.J. Garegg et al. Tetrahedron Lett. **1986**, 27, 4051-4054; B.C. Froehler Tetrahedron Lett. **1986**, 27, 5575-5578; B.C. Froehler et al. Tetrahedron Lett. **1986**, 27, 469-472; S.L. Beaucage et al. Tetrahedron Lett. **1981**, 22, 1859-1862; and U.S. Patent Nos. 5,037,882, 5,616,700, 5,571,902, 5,705,621, 5,700,922, 5,610,289, 5,635,488, 5,668,266, 5,648,480 and 5,221,736. The H-phosphonate and phosphoramidite protocols are presently preferred. More preferred is the β -cyanoethyl phosphoramidite protocol.
- 10** For the synthesis of peptides, any protocols known to those of skill in the art may be used in the methods. For example, protocols such as those described in U.S. Patent Nos. 5,726,243, 5,705,333, 4,749,742, 5,644,029, 5,637,719, 5,262,331, 4,794,150, 5,221,736 and 5,641,862 may be used.
- 15** For PNA synthesis, protocols such as those found in Brown et al. Science **1994**, 265, 777-780; Norton et al. Bioorg. & Med. Chem. **1995**, 3, 437-445; Schmidt et al. Nucl. Acids Res. **1997**, 25, 4792-4796 and U.S. Patent Nos. 5,700,922, 5,736,336, 5,719,262, 5,714,331, 5,705,333 and 5,539,083 may be used.
- 20** For oligosaccharide synthesis, protocols such as those found in U.S. Patent Nos. 5,246,840, 5,616,698, 5,374,655, 5,288,637, 5,532,147 and 5,583,042 may be used.
- All of the preceding publications and patents regarding protocols for biopolymer synthesis are incorporated by reference herein.
- 25** **3. LPCs for use in the Methods**
- LPCs for use in the methods are any of those provided herein. The LPCs have the formula $\text{Sp}(\text{X}^1)_n$, wherein Sp is a polyvalent group that has two or more, preferably three or more, more preferably 3-6, points of attachment, n corresponds to the number of points of attachment in Sp and X^1 is a reactive group that is compatible from the point of view of
- 30** biopolymer chemistry. Sp is preferably a symmetrical group such that all

X¹ groups are equivalent. It will be appreciated by those of skill in the art that such symmetry allows for uniformity in the synthesis of n biopolymers using these LPCs.

- In preferred embodiments, the LPCs used in the methods are of
- 5 formulae (I) or derivatives thereof as described above, more preferably of formulae (Ia), (Ib) and (Id), or derivatives thereof. In more preferred embodiments, the LPCs used in the methods are of formulae (IIa), (IIb) or (IIc), or derivatives thereof.

- In particularly preferred embodiments of the methods, the LPCs are
- 10 suitable for oligonucleotide synthesis. Such LPCs include the carboxyl derivatives of formulae (IIa), (IIb) or (IIc) that have the general formulae $\text{Sp}(\text{O}, \text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$ and $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$.

- In other more preferred embodiments, the LPCs used in the
- 15 methods are the carboxyl derivatives of formulae $\text{Sp}(\text{O}, \text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$ and $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$ that are further derivatized by reaction with a first monomer unit for synthesis of a biopolymer, preferably an
- 20 oligonucleotide. As exemplified herein, a first monomer unit in the synthesis of oligonucleotides may be 5'-O-dimethoxytrityl-2'-deoxythymidine or 2'-deoxythymidine. Preferred LPCs of this embodiment include tetrakis{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-dT)₄-PE-LPC), 1,3,5-tris{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenyl-methyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC), tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC), tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine
- 25
- 30

((DMT-dT)₃-Amine-LPC) and tris[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

4. Preferred Embodiments of the Method

In a preferred embodiment, the method is used to prepare
 5 oligonucleotides. This is achieved through chain elongation in the 3'→5' direction with monomeric, dimeric or trimeric phosphoramidites. The procedure represents a way to a purification oriented and effective technique for the synthesis of oligonucleotides and their analogs in preparative scale. The purification, mainly achieved by gel permeation
 10 chromatography (GPC) on Sephadex LH20, is facilitated through the coupling of the intermediates to the LPC. The considerable difference in size between reaction products and all other reagents including the excess phosphoramidites allows separation by gel filtration. The condensation product, as the largest component, is eluted first near the
 15 void volume. Moreover the LPC mediates the solubility in organic solvents during the different reaction steps and the chromatography process.

Preferably after dedimethoxytritylation the reaction products can be analyzed by MALDI-TOF mass spectrometry. Through such analysis, a direct reaction control on the step of the fully base and phosphate
 20 protected intermediates is possible. The scale of synthesis is mainly determined by the dimension of the Sephadex LH20 column and can be varied in a wide range, including up to grams or kilograms.

In an exemplary embodiment, the method involves the synthesis of the decanucleotide d(GACGGCCAGT) using monomeric phosphoramidites
 25 and an LPC based on a 1,3,5-benzene tricarboxylic acid derivative (see, Example 7).

As described in detail in Example 7, the synthesis of decanucleotide d(GACGGCCAGT) was performed using the LPC 1,3,5-tris-[9-(2'-deoxythymidine-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene
 30 ((dT)₃-Aryl-LPC). Absolute pyridine was used in the condensation reactions. 166 μmol of the LPC (corresponds to 500 μmol 5'-hydroxyl

groups) were dissolved in dry pyridine and together with a solution of 1H-tetrazole in acetonitrile added to 2.5 equivalents of the solid 5'-O-(DMT)-dG^{ib}-phosphoramidite under exclusion of moisture in an argon atmosphere. Phosphoramidite and tetrazole solution were added until no further

5 reaction was observed; a total of 2 g (2.4 mmol, 5 eq.) 5'-O-(DMT)-dG^{ib}-phosphoramidite and 12 ml (5.2 mmol, 10 eq.) tetrazole solution were used. The condensation was monitored by thin layer chromatography (TLC). The resulting mixture of product and excess reagents including several equivalents of phosphoramidite was carefully reduced to a volume

10 up to 10 ml and separated on a Sephadex LH20 column with THF/methanol 80:20 (v/v) as eluent. As expected, the condensation product eluted near the void volume, followed by the excess phosphoramidite components and finally the solvent, pyridine.

After GPC, the fractions containing the condensation product were

15 combined and oxidized with t-butylhydroperoxide (t-BuOOH) at 0 °C. The t-butylhydroperoxide (boiling point 81 °C) can easily be removed by co-evaporation with the solvents used in the oxidation (THF and methanol). The solution was reduced to dryness and the residue dissolved in dichloromethane/nitromethane/methanol 80:19:1 (v/v). For

20 dedimethoxytritylation the resulting colorless solution was mixed with trifluoroacetic acid (TFA) in the same solvent system, so that a final acid concentration not higher than 2% (by volume) was obtained. The removal of the 5'-O-protecting group was complete in less than 2 minutes and indicated by a deep red to orange color. To avoid depurination the

25 reaction was stopped by neutralization with triethylamine in an ice bath. For separation of the dedimethoxytritylated LPC compound Sephadex LH20 chromatography with a RP-silica gel filled precolumn (NUCLEOPREP® 300-30 C₁₈) was used. The precolumn was necessary to prevent contamination with dimethoxytrityl carbinol. In the first cycle the

30 intermediate d(G^{ib}T)₃-Aryl-LPC was isolated as a colorless powder after drying by co-evaporation with dioxane. The identification by MALDI-TOF

mass spectrometry showed no significant side products at that stage of synthesis.

The synthesis then followed the procedure described for the first cycle without essential changes of the method. Table 1 shows the yields, achieved in the nine cycles for condensation and oxidation (81-99%), dedimethoxytritylation (81-99%) and the whole cycle (81-99%). The total yield over all cycles amount to 33% for d(5'-O-DMT-G^{ib}A^{bz}C^{bz}G^{ib}G^{ib}C^{bz}C^{bz}A^{bz}G^{ib}T)₃-Aryl-LPC, including 26 reaction steps and 17 chromatographic separations. The quantities of reagents used in the synthesis are listed in Table 2 (see, Example 7). The used excess of β-cyanoethyl phosphoramidites varied between 5 and 12 equivalents, however, as little as 2 to 2.5 equivalents may be used. In this case the procedures for drying of the 5'-hydroxyl components were more intensive. Separation is in general quantitative. The average yield of 96% for the coupling reaction is comparable to that obtained with solid phase synthesis and better than comparable synthesis in solution. The average yield for dedimethoxytritylation is 87%. Preferred embodiments of the method avoid the use of water in chromatography. Other polar solvents for GPC on Sephadex LH20, such as pyridine or N,N-dimethylformamide, are favored for this purpose.

Table 1

Yields achieved in the oligonucleotide synthesis in solution using dT₃-Aryl-LPC for the preparation of decamer d(GACGGCCAAGT)

Synthesis of d(5'-O-DMT-G ^{ib} A ^{bz} C ^{bz} G ^{ib} G ^{ib} C ^{bz} C ^{bz} A ^{bz} G ^{ib} T) ₃ -Aryl-LPC, starting with 166 μmol dT ₃ -Aryl-LPC 1 [yields in % and μmol]:			
Cycle:	condensation:	dedimethoxytritylation:	yield per cycle:
1	87 % , 144 μmol using (DMT-dG ^{ib})-amidite	94 % , 135 μmol	81 %
2	93 % , 126 μmol using (DMT-dA ^{bz})-amidite	87 % , 110 μmol	81 %
3	91 % , 100 μmol using (DMT-dC ^{bz})-amidite	99 % , 100 μmol	91 %
4	98 % , 98 μmol using (DMT-dC ^{bz})-amidite	86 % , 84 μmol	84 %

5	5	99 % , 83 μ mol using (DMT-dG ^b)-amidite	96 % , 80 μ mol	95 %
	6	99 % , 80 μ mol using (DMT-dG ^b)-amidite	81 % , 65 μ mol	81 %
	7	99 % , 65 μ mol using (DMT-dC ^{bz})-amidite	83 % , 54 μ mol	83 %
	8	99 % , 54 μ mol using (DMT-dA ^{bz})-amidite	99 % , 54 μ mol	99 %
	9	99 % , 54 μ mol using (DMT-dG ^b)-amidite	-	-
total:		87 - 99 % , \emptyset = 96 %	81 - 99 % , \emptyset = 91 %	81 - 99 % , \emptyset = 87 %

- An inherent advantage of large scale oligonucleotide synthesis in solution in comparison to solid phase strategies is shown by the use of MALDI-TOF mass spectrometry for analysis of the fully base and phosphate protected intermediates. In contrast to solid phase synthesis, a direct control of the progressing elongation reaction is possible by analysis with this technique (for preparation of probes out of condensation, dedimethoxytritylation is necessary). For analysis of the condensation products from solid phase synthesis, cleavage of the connecting linker is necessary. The hydrolysis also leads to loss of phosphate and base protecting groups. Evidence about side reactions or incomplete reaction is therefore complicated in the analysis. The control over the basic elongation step in oligonucleotide synthesis is however of great importance for economic large scale production.

The analysis of product d(GACGGCCAGT), synthesized on dT₃-Aryl-LPC, was performed, after treatment with ammonium hydroxide, RP-HPLC and dedimethoxytritylation with acetic acid, by SMART-HPLC (Mono Q 16/5) and MALDI-TOF mass spectrometry.

In another exemplary embodiment, dimer phosphoramidites were used in the synthesis of the pentanucleotide d(GCCCT) (see, Example 8). Such building blocks are advantageous in oligonucleotide synthesis in solution because the number of coupling cycles and purification steps is reduced.

The preparation of an oligonucleotide of the sequence d(GCCCT), using the developed method for solution synthesis together with dimer phosphoramidites, demonstrated that these synthons are in general very useful. The phosphorylation is indicated by MALDI-TOF mass

- 5 spectrometry through the detected fragment ion at 3306.4 m/z, corresponding to d(5'-HO-C¹⁸C¹⁸T)₂(5'-pdT)-Aryl-LPC (theoretical mass: 3272.9 Da). The steps in one elongation cycle are performed with no significant changes to that described for monomeric phosphoramidites. Instead of 1H-tetrazole the somewhat more activating 5-(4-nitrophenyl)-
10 tetrazole is used. The coupling yield was 83%.

The following examples are included for illustrative purposes only and are not intended to limit the scope of the invention.

EXAMPLE 1

- 15 **Tetrakis-{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-dT)₄-PE-LPC)**

A. Tetrakis-[(2-cyanoethoxy)methyl]methane

- Pentaerythritol (6.84 g, 50 mmol) was dissolved in 20 ml dioxane
20 and 2 ml water and mixed up with 40 % aqueous potassium hydroxide solution (w/v, 1 ml). After cooling to 0 °C in an ice bath, acrylonitrile (16.2 g, 300 mmol) was added and the mixture stirred for 48 h at room temperature. Solvents were evaporated, the remaining oil was diluted in 100 ml dichloromethane and washed with 10 % aqueous sodium chloride
25 solution (w/v, 50 ml). After re-extraction of the aqueous layer with dichloromethane (2 x 25 ml), the combined organic layers were dried over sodium sulfate and intensively evaporated. The nitrile was obtained as a colorless syrup: yield: 14.32 g (41 mmol, 82 %). ¹H-NMR (400 MHz, CDCl₃) δ 2.60 (t, 8 H, -CH₂CN), 3.50 (s, 8 H, -C_qCH₂O-), 3.65 (t, 8 H, -
30 OCH₂-). ¹³C-NMR (100 MHz, CDCl₃) δ 18.86 (-CH₂CN), 45.68 (C_q), 65.88 (-OCH₂-), 68.82 (C_qCH₂O-), 118.22 (CN).

B. Tetrakis-[[2-(methoxycarbonyl)ethoxy]methyl]-methane

- Tetrakis-[(2-cyanoethoxy)methyl]-methane (3.48 g, 10 mmol) was dissolved in dried methanol (30 ml) and saturated with dried hydrogen chloride at room temperature. The reaction mixture was refluxed for 1 h
- 5 under continuous passing of hydrogen chloride gas. During cooling to room temperature argon was passed through the whole apparatus. After hydrolysis with ice cold water, work up was performed (150 ml) through extraction with diethyl ether (3 x 100 ml) and drying of the combined organic layers with sodium sulfate. Intensive evaporation leads to the
- 10 methyl ester as a yellowish oil: yield: 75 % (3.72 g, 7.5 mmol). ¹H-NMR (400 MHz, CDCl₃) δ 2.55 (t, 8 H, -CH₂CO-), 3.32 (s, 8 H, -C_q-CH₂O-), 3.65 (t, 8 H, -OCH₂-), 3.70 (s, 12 H, -OCH₃). ¹³C-NMR (100 MHz, CDCl₃) δ 34.92 (-CH₂CO-), 45.35 (C_q), 51.16 (CH₃), 66.60 (-OCH₂-), 69.23 (C_qCH₂O-), 172.14 (CO).

15 C. Tetrakis-(8-amino-6-aza-2-oxa-5-oxooctyl)-methane

- The methyl ester from Example I.B. (3.91 g, 8 mmol) was dissolved in dried methanol (20 ml) and mixed up with a large excess of freshly distilled ethylenediamine (dried over CaH₂) at 0 °C (200 ml, 3 mol). Under inert atmosphere (argon) the solution was stirred for 96 h at 4 °C.
- 20 Ethylenediamine was removed by evaporation and co-evaporation after dissolving in methanol (10 ml) and addition of toluene (100 ml, 3 times): yield: 88% (4.01 g, 7 mmol). ¹H-NMR (400 MHz, CDCl₃) δ 1.60 (br, 8 H, -NH₂), 2.40 (t, 8 H, -CH₂CO-), 2.82 (m, 8 H, -CH₂NH₂), 3.30 (m, 8 H, -NHCH₂-), 3.32 (s, 8 H, C_qCH₂O-), 3.68 (t, 8 H, -OCH₂-), 7.05 (m, 4 H,
- 25 NH). ¹³C-NMR (100 MHz, CDCl₃) δ 36.94 (-CH₂CO-), 41.57 (-NHCH₂-), 42.34 (CH₂NH₂), 45.37 (C_q), 67.58 (-OCH₂-), 69.73 (C_qCH₂O-), 171.90 (CO).

30 D. Tetrakis-{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}-methane ((DMT-dT)₄-PE-LPC)

The synthesis of 5'-O-dimethoxytrityl-deoxythymidine-3'-p-nitrophenylsuccinate was performed according to the procedure in

Oligonucleotide Synthesis, IRL Practical Approach Series, Hrsg.: M.J.

Gait, IRL-Press Oxford (1984). The obtained solution in dioxane/pyridine, containing the p-nitrophenyl ester (approx. 5.4 mmol) was concentrated under reduced pressure and dissolved in pyridine (10 ml). Tetrakis-(8-

- 5 amino-6-aza-2-oxa-5-oxooctyl)-methane (600 mg, 1 mmol) was added and stirred for 16 h at room temperature. Reaction mixture was concentrated and dissolved in eluent mix. The first chromatography on silica gel was performed using a step gradient from dichloromethane/ethanol 95:5 (v/v) to dichloromethane/ethanol 60:40 (v/v). All solvents contained 0.5% pyridine (v/v). Further purification with gel permeation chromatography on Sephadex LH20 gave only small portions of pure material (eluent: THF/isopropanol 80:20, v/v). Repeated silica gel column chromatography (with dichloromethane/ethanol, 15 to 30% ethanol, 0.5% pyridine) and gel permeation chromatography gave the product in colorless crystalline form, total yield: 45% (1.4 g, 0.45 mmol). ¹H-NMR (400 MHz, CDCl₃) δ 1.37 (s, 12 H, CH₃-, Base), 2.40 (m, 8 H, -CH₂CO-), 2.40/2.50 (m, 8 H, H2'), 2.49 (m, 8 H, -COCH₂-), 2.63 (m, 8 H, -CH₂CO-), 3.29 (s, 8 H, C_qCH₂O), 3.36 (m, 16 H, 2 x -NHCH₂-) 3.43 (m, 8 H, H5'), 3.68 (m, 8 H, -OCH₂-), 3.76 (s, 24 H, CH₃O-), 4.15 (m, 4 H, H4'), 5.43 (m, 4 H, H3'), 6.35 (dd, 4 H, H1'), 6.8 - 7.4 (m, 52 H, H_{Ar}), 7.60 (s, 4 H, CH, Base), 8.62 (dd, 2 x 4 H, 2 x -NH-CH₂-), 9.90 (s, 4 H, NH, Base). ¹³C-NMR (100 MHz, CDCl₃) δ 11.63 (CH₃, Base), 29.74 (C12), 30.56 (C11), 36.79 (C4), 37.80 (C2'), 39.38 (C8), 40.01 (C7), 55.28 (-OCH₃), 63.85 (C5'), 67.50 (C3), 69.24 (C1), 75.88 (C3'), 83.81/84.30 (C1'/C4'), 87.22 (C₂, DMT), 111.93 (C5, Base), 113.34 (C3, DMT), 127.26/128.06/128.13 (C2'/C3'/C4', DMT), 130.09 (C2, DMT), 135.09 (C1, DMT), 135.40 (C6, Base), 144.16 (C1', DMT), 151.34 (C2, Base), 158.75 (C4, DMT), 164.24 (C4, Base), 171.96 (C5 and C10), 172.16 (C13).
- 25

EXAMPLE 2

1,3,5-Tris-{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC)

5 A. 1,3,5-Benzenetricarboxylic acid tris-N-(2-aminoethyl)amide

1,3,5-Benzenetricarboxylic acid trimethyl ester (1.04 g 4.0 mmol) was treated with ethylenediamine (66.7 ml, 1 mol), following the procedure described in Example 1.C. The reaction was complete after 24 h. After lyophilization with dioxane the product was obtained as a white solid; yield: 99 % (1.35 g, 4.0 mmol). ¹H-NMR (400 MHz, d₆-DMSO) δ 1.55 (br, 6 H, -NH₂), 2.72 (m, 6 H, -CH₂NH₂), 3.32 (m, 6 H, -NHCH₂-), 8.42 (s, 3 H, CH_{Ar}), 8.65 (t, 3 H, NH). ¹³C-NMR (100 MHz, d₆-DMSO) δ 42.16 (-CH₂NH₂), 44.16 (-NHCH₂-), 129.30 (CH_{Ar}), 135.95 (C_{Ar}), 166.56 (CO).

15 B. 1,3,5-Tris-{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC)

A solution of 5'-O-dimethoxytrityl-deoxythymidine-3'-p-nitrophenylsuccinate (approx. 10 mmol) in dioxane/pyridine was concentrated under reduced pressure and dissolved in DMF (25 ml) and pyridine (10 ml). 1,3,5-Benzenetricarboxylic acid tris-N-(2-aminoethyl)amide (Example 2.B.) (673 mg, 2 mmol) was added, the reaction mixture stirred for 16 h at room temperature, concentrated and dissolved in eluent mix. Silica gel column chromatography (step gradient with dichloromethane and 2 to 20% ethanol, 0.5% pyridine, v/v) led to mostly pure product fractions. Gel permeation chromatography on Sephadex LH20 (eluent: THF/methanol 80:20 v/v) of the impure fractions gave additional product, total yield: 90 % (4.0 g, 1.8 mmol). ¹H-NMR (400 MHz, CDCl₃) δ 1.32 (s, 9 H, CH₃-, Base), 2.40/2.50 (m, 6 H, H2'), 2.44 (m, 6 H, -COCH₂-), 2.50 (m, 6 H, -CH₂CO-), 3.40 (m, 6 H, H5'), 3.54 (m, 12 H, 2 x -NHCH₂-), 3.77 (s, 18 H, CH₃O-), 4.15 (m, 3 H, H4'), 5.40 (m, 3 H, H3'), 6.32 (dd, 3 H, H1'), 6.8 - 7.4 (m, 39 H, H_{Ar}), 7.60 (s,

- 3 H, CH, Base), 8.19 (dd, 2 x 3 H, 2 x -NH-CH₂-), 8.40 (s, 3 H, CH_{Ar}).
¹³C-NMR (100 MHz, CDCl₃) δ 11.66 (CH₃, Base), 29.97 (C8), 30.66 (C7), 37.57 (C2'), 39.38 (C4), 40.57 (C3), 55.26 (-OCH₃), 63.79 (C5'), 75.80 (C3'), 83.58/84.27 (C1'/C4'), 87.21 (C₂, DMT), 112.06 (C5, Base),
 5 113.34 (C3, DMT), 127.24/128.13/128.23 (C2'/C3'/C4', DMT), 129.04 (CH_{Ar}), 130.10 (C2, DMT), 134.93 (C6, Base), 135.16 (C1, DMT), 135.27 (C_{Ar}), 144.21 (C1', DMT), 151.34 (C2, Base), 158.79 (C4, DMT), 164.08 (C4, Base), 166.57 (C1), 171.95 (C6), 172.16 (C9).

EXAMPLE 3

10 Tetrakis-[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC)

- Fractogel[®] EMD-SO₃⁻ 650 (M) (Merck, Darmstadt, Germany) in protonated form (counterion H⁺) was used for dedimethoxytritylation and solid phase extraction. (DMT-dT)₄-PE-LPC (Example 1) (466 mg, 150
 15 μmol) was dissolved in THF/ethanol 95:5 (v/v) and transferred to a column (inner diameter: 20 mm) containing 50 ml ion exchanger, equilibrated with n-hexane/THF 50:50 (v/v). Dimethoxytrityl-compounds were washed off with n-hexane/THF/ethanol 50:50:5 (v/v, 100 ml), THF/ethanol 95:5 (v/v, 100 ml) and THF/Ethanol 50:50 (v/v, 150 ml).
 20 dT₄-PE-LPC was eluted with pure ethanol, solvent was removed and thoroughly co-evaporation with toluene gave the product as a colorless solid, yield: 93 % (263 mg, 139 μmol). ¹H-NMR (d₆-DMSO) δ 1.80 (s, 16 H, CH₃-, Base), 2.20 (m, 8 H, H2'), 2.25 (m, 8 H, -CH₂CO-), 2.38 (m, 8 H, -COCH₂-), 2.52 (m, 8 H, -CH₂CO-), 3.08 (m, 16 H, 2 x -NHCH₂-), 3.21
 25 (s, 8 H, C_qCH₂O), 3.51 (m, 8 H, -OCH₂-), 3.64 (m, 8 H, H5'), 3.98 (m, 4 H, H4'), 5.22 (m, 4 H, H3'), 6.18 (dd, 4 H, H1'), 7.75 (s, 4 H, CH, Base), 7.80/7.85 (dd, 2 x 4 H, 2 x -NH-CH₂-). ¹³C-NMR (d₆-DMSO) δ 12.18 (CH₃, Base), 29.02 (C12), 29.94 (C11), 36.06 (C4), 36.41 (C2'), 38.35 (C8), 38.84 (C7), 44.70 (C_q), 61.27 (C5'), 67.19 (C3), 69.00 (C1),
 30 74.68 (C3'), 83.64/84.50 (C1'/C4'), 109.64 (C5, Base), 135.73 (C6,

Base), 150.39 (C2, Base), 163.58 (C4, Base), 170.30 (C5), 170.75 (C10), 171.94 (C13).

EXAMPLE 4

5 1,3,5-Tris-[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC)

Procedure 1

Dedimethoxytritylation and solid phase extraction with Fractogel[®] EMD-SO₃⁻ 650 (M) (counterion H⁺), additional purification with gel permeation chromatography using Sephadex LH20: (DMT-dT)₃-Aryl-LPC
 10 (Example 2) (1.1 g, 500 μmol) was dissolved in THF/ethanol 99:1 (v/v) and transferred to a column (inner diameter: 30 mm), containing 150 ml ion exchanger, equilibrated with n-hexane/THF/ethanol 60:35:5 (v/v). Dimethoxytrityl-compounds were washed off with n-hexane/THF/ethanol 60:35:5 (v/v, 200 ml), THF/n-hexane/ethanol 60:35:5 (v/v, 200 ml) and
 15 THF/ethanol 95:5 (v/v, 200 ml). dT₃-Aryl-LPC was eluted with THF/ethanol 65:35 and THF/ethanol 50:50 (v/v, 300 ml each). The crude product was purified with gel permeation chromatography on Sephadex LH20 (eluent: THF/isopropanol 80:20 v/v), giving a colorless solid after removal of solvents, yield 82% (530 mg, 410 μmol).

20 Procedure 2

Dedimethoxytritylation with TFA reagent, purification with solid phase extraction using LiChroprep[®]-DIOL (Merck): (DMT-T)₃-Aryl-LPC (Example 2) (580 mg, 260 μmol) was dissolved in (10 ml) 1,2-dichloroethane/nitromethane/methanol 80:19:1 (v/v) and TFA solution in the
 25 same solvent system was added (3%, 7.85 mmol TFA, 20 ml). After 3 minutes the reaction mixture was transferred to a column (inner diameter: 25 mm) filled with LiChroprep[®]-DIOL (90 ml), equilibrated with dichloromethane/ethanol 95:5 (v/v). The same solvent mixture was used for elution of dimethoxytrityl compounds (300 ml). Washing with
 30 dichloromethane/pyridine/ethanol 50:40:10 (v/v) gave the product dT₃-

Aryl-LPC after removal of solvents and co-evaporation with toluene (2 x 20 ml) as a colorless crystalline solid, yield: 97 % (330 mg, 252 μ mol).

Procedure 3

- Dedimethoxytritylation with TFA reagent, purification by gel permeation chromatography using Sephadex LH20 and additional precipitation in diethyl ether: A TFA solution in 1,2-dichloroethane/nitromethane/methanol 80:19:1 (v/v, 2% TFA, 10 mmol) was dropwise mixed with a solution of (DMT-dT)₃-Aryl-LPC (Example 2) (739 mg, 330 μ mol) in the same solvent system (5 ml) under stirring at room temperature. After 2 minutes the reaction mixture was cooled in an ice bath and triethylamine (10 mmol in 4 ml of the solvent system) was added. Solvents were removed under reduced pressure and the residue dissolved in N,N-dimethylformamide (5 ml). After chromatography on Sephadex LH20 (eluent: THF/water 70:30, v/v) the most product containing fractions were contaminated with traces of dimethoxytrityl compounds. After removal of the solvents and co-evaporation with dioxane (40 ml), the residue was dissolved in THF/methanol (10 ml) and precipitated into heavily stirred diethyl ether (300 ml). Decantation of the ether layer, filtration of the precipitate under washing with diethyl ether and co-evaporation with pyridine (5 ml) and toluene (2 x 10 ml) gave dT₃-Aryl-LPC as a colorless solid, yield: 88% (380 mg, 290 μ mol).

Procedure 4

- Dedimethoxytritylation with TFA reagent and neutralization with trioctylamine, purification by chromatography using NUCLEOPREP 300-30 C₁₈: (DMT-dT)₃-Aryl-LPC (Example 2) (113 mg, 50 μ mol) was dissolved in 1,2-dichloroethane/nitro-methane/methanol 80:19:1 (v/v, 4 ml) and a TFA solution in the same solvent system (1.5 mmol TFA in 1 ml) was added dropwise. After 3 minutes the reaction mixture was cooled in an ice bath and neutralized by trioctylamine (1.5 mmol in 1 ml of the solvent system). Solvents were removed under reduced pressure and the residue dissolved in acetonitrile/THF/water 56:14:30 (v/v, 4 ml), also the eluent in the

- following chromatography using NUCLEOPREP 300-30 C18 (column: 310 x 20 mm, flow 10 ml/min). Product is present in fractions with a retention time between 7 and 9 minutes and well separated of all other components of the probe. After removal of solvents under reduced
- 5 pressure and lyophilization with dioxane dT₃-Aryl-LPC is obtained as a colorless solid, yield: 98 % (64 mg, 49 μmol).

Procedure 5

- Dedimethoxytritylation with TFA reagent, purification by combined chromatography using NUCLEOPREP 300-30 C18 and Sephadex LH20:
- 10 (DMT-dT)₃-Aryl-LPC (Example 2) (850 mg, 384 μmol) was dissolved in 1,2-dichloroethane/nitro-methane/methanol 80:19:1 (v/v, 33 ml) and mixed with a TFA solution in the same solvent mixture (11.5 mmol TFA in 10 ml). After 2 minutes a triethylamine solution was added for neutralization under cooling in an ice bath (11.5 mmol, 5 ml in the same
- 15 solvent system). The reaction mixture was concentrated under reduced pressure and dissolved with the eluent THF/water 60:40 (v/v, 5 ml). Chromatography was performed with a precolumn filled with NUCLEOPREP 300-30 C18 (310 x 20 mm) and a second column with Sephadex LH20 (460 x 30 mm, flow: 1 ml/min). Product containing
- 20 fractions were combined, concentrated and co-evaporated with dioxane (2 x 40 ml). Final lyophilization with dioxane (10 ml) gave dT₃-Aryl-LPC as a colorless solid, yield: 97% (490 mg, 374 μmol).
- ¹H-NMR (400 MHz, d₆-DMSO) δ 1.78 (s, 9 H, CH₃-, Base), 2.25 (m, 6 H, H2'), 2.40 (m, 6 H, -COCH₂-), 2.55 (m, 6 H, -CH₂CO-), 3.24/3.34 (m, 12
- 25 H, 2 x -NHCH₂-), 3.60 (m, 6 H, H5'), 3.97 (m, 3 H, H4'), 5.20 (m, 3 H, H3'), 6.17 (dd, 3 H, H1'), 7.70 (s, 3 H, CH, Base), 8.00 (dd, 3 H, -NHCH₂-), 8.40 (s, 3 H, CH_{Ar}), 8.68 (dd, 3 H, -NHCH₂). ¹³C-NMR (100 MHz, d₆-DMSO) δ 12.19 (CH₃, Base), 29.02 (C8), 29.76 (C7), 36.38 (C2'), 38.18 (C4), 39.13 (C3), 61.26 (C5'), 74.68 (C3'), 83.60/84.47
- 30 (C1'/C4'), 109.64 (C5, Base), 128.48 (CH_{Ar}), 134.78 (C_{Ar}), 135.74 (C6,

Base), 150.39 (C2, Base), 163.57 (C4, Base), 165.51 (C1), 170.79 (C6), 171.95 (C9).

EXAMPLE 5

5 Tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine ((DMT-dT)₃-Amine-LPC)

To a solution of 5'-O-dimethoxytrityl-deoxythymidine-3'-p-nitrophenylsuccinate (approx. 9.4 mmol) in dioxane/pyridine a solution of tris-(2-aminoethyl)-amine (292 mg, 2 mmol) in dried acetonitrile (10 ml) was added and stirred for 12 h at room temperature. The reaction mixture

10 was concentrated and the residue dissolved in dichloromethane (containing 0.5% pyridine). Silica gel column chromatography was started with dichloromethane/methanol (0.5% pyridine), increasing the methanol content stepwise up to 15%. Product containing fractions were combined, concentrated and precipitated into n-hexane/diethyl ether 3:1

15 (v/v). (DMT-dT)₃-Amine-LPC was isolated by filtration as a colorless solid, yield: 74 % (2.99 g, 1.48 mmol). ¹H-NMR (CDCl₃) δ 1.35 (s, 9 H, CH₃-, Base), 2.40/2.50 (m, 6 H, H2'), 2.50/2.52/ 2.62 (m, 3 x 6 H, -COCH₂-, -CH₂N_{central}-, -CH₂CO-), 3.23 (m, 6 H, -CONHCH₂-), 3.45 (m, 6 H, H5'), 3.76 (s, 18 H, CH₃O-), 4.15 (m, 3 H, H4'), 5.40 (m, 3 H, H3'), 6.38 (dd,

20 3 H, H1'), 6.8 - 7.4 (m, 39 H, H_{Ar}), 7.58 (CH, Base), 9.55 (NH, Base). ¹³C-NMR (CDCl₃) δ 11.60 (CH₃, Base), 29.77 (C6), 30.46 (C5), 37.75/37.75 (C2'/C2), 54.63 (C1), 55.26 (-OCH₃), 63.88 (C5'), 75.89 (C3'), 83.84/84.39 (C1'/C4'), 87.19 (C_z, DMT), 111.89 (C5, Base), 113.35 (C3, DMT), 127.21/128.04/128.23 (C2'/C3'/C4', DMT), 130.08

25 (C2, DMT), 135.18 (C1, DMT), 135.39 (C6, Base), 144.24 (C1', DMT), 150.91 (C2, Base), 158.78 (C4, DMT), 163.87 (C4, Base), 171.77 (C4), 172.25 (C7).

EXAMPLE 6

Tris-[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC)

- Dedimethoxytritylation of (DMT-dT)₃-Amine-LPC (Example 5) (660 mg, 325 μ mol) was performed with TFA (2%) in nitromethane/methanol (96:2, v/v, 40 ml). Purification was achieved by solid phase extraction using LiChroprep®-NH₂ (Merck, 50 ml, column: 150 x 20 mm). Dimethoxytrityl compounds were eluted with n-hexane/ethyl acetate 2:1 (v/v, 50 ml), the nucleotidic compounds with ethyl acetate/ethanol 2:1 (v/v, 150 ml). Product containing fractions were concentrated and gel permeation chromatography was used for further purification (Sephadex LH60, eluent: THF/ethanol 90:10, v/v; column: 310 x 25 mm, flow 1 ml/min). dT₃-Amine-LPC was isolated as a colorless solid, yield: 61% (220 mg, 197 μ mol). ¹H-NMR (d₆-DMSO) δ 1.77 (s, 9 H, CH₃-, Base), 2.25 (m, 6 H, H2'), 2.40 (m, 6 H, -COCH₂-), 2.48 (m, 6 H, -CH₂N_{central}), 2.55 (m, 6 H, -CH₂CO), 3.08 (m, 6 H, -CONHCH₂-), 3.62 (m, 6 H, H5'), 3.98 (m, 3 H, H4'), 5.18 (m, 3 H, H3'), 6.18 (dd, 3 H, H1'), 7.72 (CH, Base), 7.75 (dd, 3 H, -NHCH₂-). ¹³C-NMR (d₆-DMSO) δ 12.18 (CH₃, Base), 29.06 (C6), 29.70 (C5), 36.43 (C2'), 36.98 (C2), 53.35 (C1), 61.27 (C5'), 74.67 (C3'), 83.63/84.51 (C1'/C4'), 109.63 (C5, Base), 135.72 (C6, Base), 150.38 (C2, Base), 163.57 (C4, Base), 170.59 (C4), 171.94 (C7).

EXAMPLE 7

Synthesis of d(5'-O-DMT-G^{ib}pA^{bz}pC^{bz}pG^{ib}pG^{ib}pC^{bz}pC^{bz}pA^{bz}pG^{ib}pT)₃-Aryl-LPC

- The 5'-hydroxyl component dT₃-Aryl-LPC (Example 4) (228 mg 166 μ mol) was co-evaporated with dried pyridine (3 x 10 ml) and dissolved in the same solvent (10 ml). (DMT)-dG^{ib}-phosphoramidite (1.0 g, 1.2 mmol) was submitted in a 100 ml-two-necked reaction flask under argon atmosphere. The solution of the 5'-hydroxyl component and tetrazole (6 ml, 30.8 mg/ml in acetonitrile, 2.6 mmol) were dropped in 500 μ l portions via syringe through septum to phosphoramidite. In case of incomplete reaction phosphoramidite as solid and tetrazole solution were added

(controlled by TLC). After complete reaction the mixture was concentrated under reduced pressure and condensation products isolated by GPC using Sephadex LH20 (eluent: THF/methanol 80:20 v/v, column: 530 x 30 mm, flow: 1 ml/min) without oxidation. Product containing

5 fractions were combined, reduced to a volume of 10 ml and oxidized at 0 °C with t-butylhydroperoxide (80% solution in di-t-butylhydroperoxide, 200 μ l, 1.5 mmol) for 5 min. After concentrating and intensive drying in vacuo (5'-O-DMT-G^{ib}pT)₃-Aryl-LPC was obtained as a colorless solid, yield 87% (520 mg, 144 μ mol).

- 10 The compound was dedimethoxytritylated with TFA reagent, using dichloromethane instead of 1,2-dichloroethane, following the procedure described in Example 4. The amounts of used reagents are given in Table 2, the chromatographic conditions were the same. The intermediate d(G^{ib}pT)₃-Aryl-LPC was analyzed via MALDI-TOF mass spectrometry
- 15 (M + H⁺: theoretical mass 2666.39 Da, found: 2678.3 m/z).

According to the above described cycle, the further elongation steps for the synthesis of d(5'-O-DMT-

G^{ib}pA^{bz}pC^{bz}pG^{ib}pG^{ib}pC^{bz}pC^{bz}pA^{bz}pG^{ib}pT)₃-Aryl-LPC were performed, using the amounts of reagents listed in Table 2. Data of analysis of the

- 20 intermediates via MALDI-TOF mass spectrometry are listed in Table 3. Over all nine cycles the crude compound d(5'-O-DMT-

G^{ib}pA^{bz}pC^{bz}pG^{ib}pG^{ib}pC^{bz}pC^{bz}pA^{bz}pG^{ib}pT)₃-Aryl-LPC was obtained in a yield of 33% (54 μ mol, 750 mg).

Table 2

- 25 The amounts of reagents used in the different reaction steps and cycles in solution synthesis of d(5'-O-DMT-G^{ib}pA^{bz}pC^{bz}pG^{ib}pG^{ib}pC^{bz}pC^{bz}pA^{bz}pG^{ib}pT)₃-Aryl-LPC

	5'-hydroxyl-component:	phosphor-amidite:	tetrazole:	t-BuOOH:	yield:	dedimethoxy-trityl.:TFA/Et ₃ N	yield:
30 1.cycle 81 %	166 μ mol in 10 ml pyridine	2 g (dG ^{ib}) 2.4 mmol 5 eq.	12 ml 5.2 mmol 10 eq.	200 μ l 1.5 mmol 3 eq.	520 mg 144 μ mol 87 %	1.14/2.1 ml 15 mmol 30 eq.	360 mg 135 μ mol 94 %
2.cycle 81 %	135 μ mol in 10 ml pyridine	3 g (dA ^{bz}) 3.5 mmol 8.75 equiv.	18 ml 7.8 mmol 19 equiv.	200 μ l 1.5 mmol 3.7 equiv.	630 mg 126 μ mol 93 %	1.14/2.1 ml 15 mmol 30 equiv.	450 mg 110 μ mol 87 %

	5'-hydroxyl-component:	phosphor-amidite:	tetrazole:	t-BuOOH:	yield:	dedimethoxytrityl.:TFA/Et ₃ N	yield:
5	3.cycle 91 %	110 μ mol in 10 ml pyridine	1.5g (dC ^{bz}) 1.8 mmol 5.5 equiv.	12 ml 5.2 mmol 16 equiv.	200 μ l 1.5 mmol 4.5 equiv.	630 mg 100 μ mol 91 %	0.57/1.1 ml 7.5 mmol 23 equiv. 580 mg 100 μ mol 99 %
	4.cycle 84 %	100 μ mol in 10 ml pyridine	2 g (dC ^{bz}) 2.4 mmol 8 equiv.	12 ml 5.2 mmol 17 equiv.	200 μ l 1.5 mmol 5 equiv.	750 mg 98 μ mol 98 %	0.57/1.1 ml 7.5 mmol 25 equiv. 570 mg 84 μ mol 86 %
	5.cycle 95 %	84 μ mol in 10 ml pyridine	1.5g (dG ^{ib}) 1.8 mmol 7 equiv.	9 ml 3.9 mmol 15 equiv.	200 μ l 1.5 mmol 6 equiv.	750 mg 83 μ mol 99 %	0.57/1.1 ml 7.5 mmol 30 equiv. 650 mg 80 μ mol 96 %
	6.cycle 81 %	80 μ mol in 10 ml pyridine	1.5g (dG ^{ib}) 1.8 mmol 7.5 equiv.	9 ml 3.9 mmol 16 equiv.	200 μ l 1.5 mmol 6 equiv.	850 mg 80 μ mol 99 %	0.57/1.1 ml 7.5 mmol 31 equiv. 620 mg 65 μ mol 81 %
10	7.cycle 83 %	65 μ mol in 10 ml pyridine	2 g (dC ^{bz}) 2.4 mmol 12 equiv.	12 ml 5.2 mmol 26 equiv.	200 μ l 1.5 mmol 7 equiv.	770 mg 65 μ mol 99 %	0.57/1.1 ml 7.5 mmol 38 equiv. 580 mg 54 μ mol 83 %
	8.cycle 99 %	54 μ mol in 10 ml pyridine	1.4g (dA ^{bz}) 1.6 mmol 10 equiv.	9 ml 3.9 mmol 24 equiv.	200 μ l 1.5 mmol 10 equiv.	750 mg 54 μ mol 99 %	0.57/1.1 ml 7.5 mmol 46 equiv. 660 mg 54 μ mol 99 %
	9.cycle	54 μ mol in 10 ml pyridine	2 g (dG ^{ib}) 2.4 mmol 15 equiv.	12 ml 5.2 mmol 30 equiv.	200 μ l 1.5 mmol 10 equiv.	750 mg 54 μ mol 99 %	-

- 15 In Table 3 the MALDI-TOF mass spectrometry data, achieved from monitoring the complete synthesis are listed. The dedimethoxytritylated intermediates were dissolved in THF/water 70:30 (v/v) and mixed with matrix solution in the same solvent (3-hydroxypicolinic acid, 0.7 mol/l). The 5'-hydroxyl components could be satisfactory identified up to cycle
- 20 6. Deviations from the calculated mass and mass differences are observed at the last three cycles, probably through the increasing occurrence of side products in synthesis and fragmentation during desorption and ionization, leading to signals, which are strongly broadened.

25

Table 3

Data achieved from 5'-hydroxyl components via MALDI-TOF mass spectrometry.

intermediates analyzed after dedimethoxytritylation	theoretical mass [Da]	found: M + H ⁺ (m/z)	introduced mass difference
30 5'-hydroxyl component in 1. cycle: d(T) ₃ -Aryl-LPC	1309.26	1325.9	
5'-hydroxyl component in 2. cycle: d(G ^b pT) ₃ -Aryl-LPC	2666.39	2678.3 $\Delta m = 1352$	+ (3xdG ^b p) $\Delta m = 1357$

5	5'-hydroxyl component in 3. cycle: d(A ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	4077.49	4093.7 Δm = 1415	+ (3xdA ^{bz} p) Δm = 1411
	5'-hydroxyl component in 4. cycle: d(C ^{bz} pA ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	5416.54	5426.1 Δm = 1332	+ (3xdC ^{bz} p) Δm = 1339
	5'-hydroxyl component in 5. cycle: d(C ^{bz} pC ^{bz} pA ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	6755.59	6764.6 Δm = 1338	+ (3xdC ^{bz} p) Δm = 1339
	5'-hydroxyl component in 6. cycle: d(G ^{ib} pC ^{bz} pC ^{bz} pA ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	8112.83	8112.8 Δm = 1366	+ (3xdG ^{ib} p) Δm = 1357
10	5'-hydroxyl component in 7. cycle: d(G ^{ib} pG ^{ib} pC ^{bz} pC ^{bz} pA ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	9469.11	9346.8 Δm = 1234	+ (3xdG ^{ib} p) Δm = 1357
	5'-hydroxyl component in 8. cycle: dC ^{bz} pG ^{ib} pG ^{ib} pC ^{bz} pC ^{bz} pA ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	10808.96	10774.8 Δm = 1428	+ (3xdC ^{bz} p) Δm = 1339
15	5'-hydroxyl component in 9. cycle: d(A ^{bz} pC ^{bz} pG ^{ib} pG ^{ib} pC ^{bz} pC ^{bz} pA ^{bz} pG ^{ib} pT) ₃ -Aryl-LPC	12219.97	12143.2 Δm = 1368	+ (3xdA ^{bz} p) Δm = 1411

- For cleavage of decamer d(GACCGGCAGT) from the liquid phase carrier and for base and phosphate deprotection, the compound d(5'-O-DMT-G^{ib}pA^{bz}pC^{bz}pG^{ib}pG^{ib}pC^{bz}pC^{bz}pA^{bz}pG^{ib}pT)₃-Aryl-LPC (13 mg, 1 μmol) was treated with ammonium hydroxide (32%, 3 ml) for 20 h at 55 °C.
- 20 The characterization of main and side products starts with reversed phase HPLC using the DMT group for separation. 200 μmol were purified using a Waters chromatography system Delta Prep 4000 (column: Whatman, Partisil 10 ODS M9, 500 x 9.4 mm, 10 μm particles, eluents were: 0.1 M tri-ethylammonium acetate at pH 7.0 (A) and acetonitrile (B), flow: 4
- 25 ml/min, 5 to 40 % B in 40 min, monitored at 260 nm). Fractions were collected around the peak maximum at 32 min, followed by dedimethoxytritylation with acetic acid (80%, 500 μl). The product was analyzed by ion exchange HPLC using the Pharmacia SMART system (column: Mono Q 1.6/5, eluents were 25 mM tris-HCl, 1 mM EDTA, pH
- 30 8.0 (A) and: 25 mM Tris-HCl, 1 mM EDTA, 1 M NaCl, pH 8.0 (B), flow: 100 μl/min: start: 20% B, 4 min isocratic, 20 to 60% B in 26 min) and compared to d(GACCGGCAGT), achieved from solid phase synthesis on CPG with included capping step.

EXAMPLE 8

Synthesis with dimer phosphoramidites

- The 5'-hydroxyl component dT₃-Aryl-LPC (Example 4) (131 mg, 100 μ mol, in 5 ml pyridine) was treated with d(5'-O-DMT-C^{tl}pC^{tl})
- 5 phosphoramidite (total: 1.79 g, 1.36 mmol, 4.5 equivalents) in the above described procedures. Instead of 1H-tetrazole, 5-(4-nitrophenyl)-tetrazole was used (1.4 mmol, as suspension in acetonitrile). Oxidation with t-butylhydroperoxide (80% in di-t-butylperoxide, 3.75 mmol, 500 μ l, at 0 °C) was performed before GPC with Sephadex LH20 (Eluent: THF/methanol 80:20, same conditions as described above). Yield: 99% crude (5'-O-DMT-C^{tl}pC^{tl}pT)₃-Aryl-LPC (516 mg, 99 μ mol). In the dedimethoxy-tritylation step, TFA (760 μ l) in dichloromethane/nitromethane/methanol 80:19:1 (32 ml, v/v). and triethylamine (1.4 ml at 0 °C) was used. Purification was obtained by the described chromatography using the
- 15 eluent THF/water 60:40 (v/v) and the precolumn filled with NUCLEOPREP 300-30 C18 (310 x 20 mm) combined with the second column with Sephadex LH20 (460 x 30 mm, flow: 1 ml/min). Yield per cycle: 83% (340 mg, 83 μ mol d(C^{tl}pC^{tl}pT)₃-Aryl-LPC). MALDI-TOF-MS: (MH⁺ = 4108.1 m/z, M_{calc.} = 4077.56). In the second condensation 4.4
- 20 equivalents of d(5'-O-DMT-G^{DPC/lb}pC^{tl}-phosphoramidite (1.1 mmol, 1.65 g) and 5-(4-nitrophenyl)-tetrazole (1.2 mmol) were used leading to compound d(5'-O-DMT-G^{DPC/lb}pC^{tl}pC^{tl}pC^{tl}pT)₃-Aryl-LPC (570 mg, 69 μ mol, 83%). Analysis was performed as described above, but varying the eluent conditions in SMART-HPLC (Start: 0% B, 4 min isocratic, 0 to 60% B in
- 25 26 min). MALDI-TOF-MS for d(GpCpCpCpT): M + H⁺: 1440.5 m/z (M_{calc.} = 1435.0 Da), d(GpCpCpCp): M + H⁺: 1194.8 m/z (M_{calc.} = 1192.8 Da).

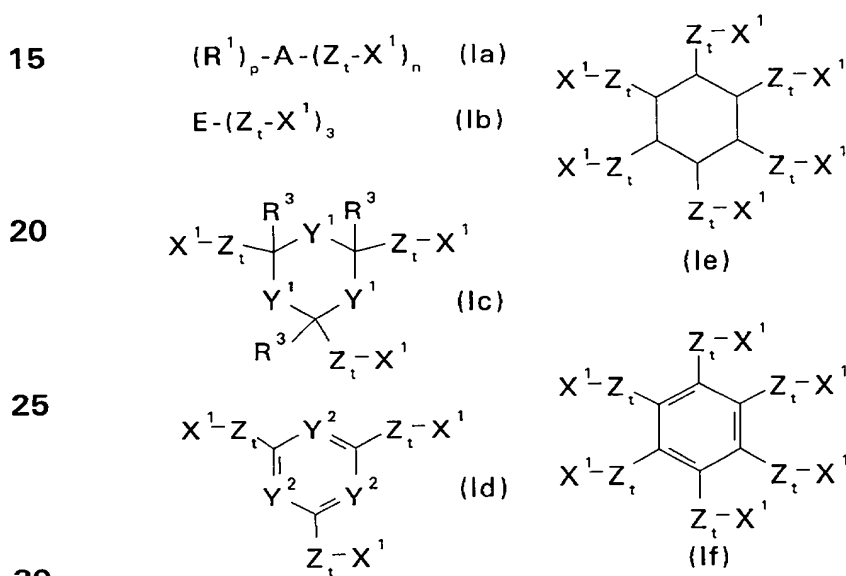
Since modifications will be apparent to those of skill in this art, it is intended that this invention be limited only by the scope of the appended

30 claims.

WHAT IS CLAIMED IS:

1. A liquid phase carrier (LPC) of formula $\text{Sp}(\text{X}^1)_n$, wherein: Sp is a polyvalent group that has more than two points of attachment, n is the number of points of attachment in Sp and X^1 is a reactive group for
5 synthesis of biopolymers.
2. The LPC of claim 1, wherein: Sp is a symmetrical group such that all X^1 groups are equivalent.
3. The LPC of claim 1, wherein n is 3-6.
4. The LPC of claim 1, wherein: X^1 is OH, SH, NH_2 , COR^5 or
10 COOR^4 , where R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclalkyl; and R^5 is halide, heteroaryl or pseudohalide.

5. The LPC of claim 1 that has formulae (I):



- wherein: A is carbon or silicon; E is nitrogen or P(O); R^1 and R^3 are each independently hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocyclalkyl; p is 0 or 1; Z is any combination of 1-12
35 units selected from 1,2-, 1,3- or 1,4-phenylene and alkylene units, which units may be combined in any order, with the proviso that if the LPC is of formula (Ia) or (Ib), then Z contains at least two phenylene or methylene

units; t is 1; X^1 is any reactive group which can be used in biopolymer synthesis; n is 3 or 4; Y^1 is CH_2 , NH, S or O; Y^2 is selected from CH and N; R^1 , R^3 , X^1 , Y^1 , Y^2 and Z are unsubstituted or substituted with one or more substituents each independently selected from Q; and Q is halogen,

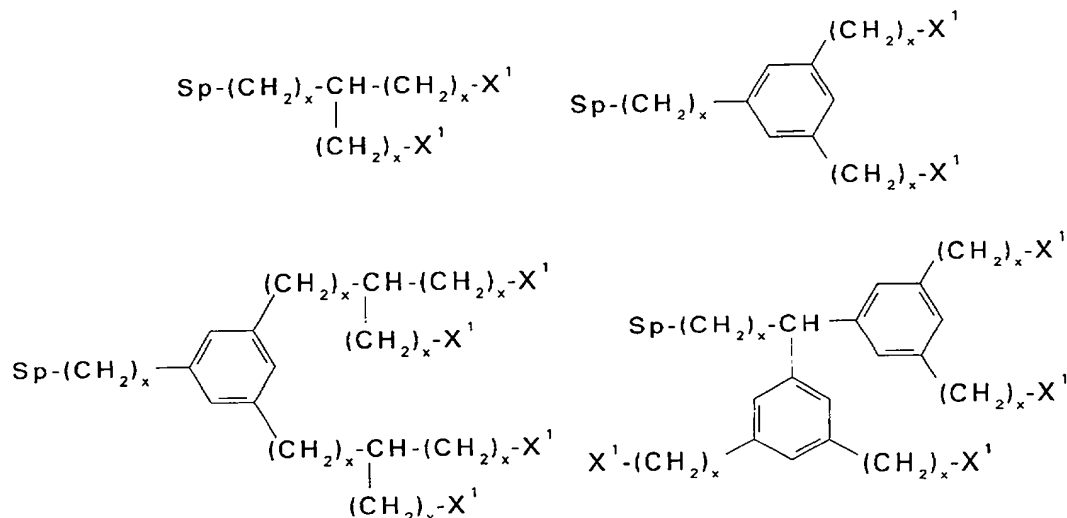
- 5 hydroxy, nitrile, nitro, formyl, mercapto, carboxy, alkyl, haloalkyl, polyhaloalkyl, aminoalkyl, diaminoalkyl, alkenyl containing 1 to 2 double bonds, alkynyl containing 1 to 2 triple bonds, cycloalkyl, cycloalkylalkyl, aryl, heteroaryl, arylalkyl, heteroarylalkyl, alkylidene, arylalkylidene, alkylcarbonyl, arylcarbonyl, heteroarylcarbonyl, alkoxycarbonyl, alkoxycarbonylalkyl, aryloxycarbonyl, aryloxycarbonylalkyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, arylaminocarbonyl, diarylaminocarbonyl, arylalkylaminocarbonyl, alkoxy, aryloxy, perfluoroalkoxy, alkenyloxy, alkynyloxy, arylalkoxy, amino, aminoalkyl, alkylaminoalkyl, dialkylaminoalkyl, arylaminoalkyl, diarylaminoalkyl, alkylamino, dialkylamino, arylamino, diarylamino, alkylaryl amino, alkylcarbonylamino, alkoxycarbonylamino, arylcarbonylamino, aryloxycarbonylamino, azido, alkylthio, arylthio, perfluoroalkylthio, thiocyano, isothiocyano, alkylsulfinyl, alkylsulfonyl, arylsulfinyl, arylsulfonyl, aminosulfonyl, alkylaminosulfonyl, dialkylaminosulfonyl, arylaminosulfonyl or diarylaminosulfonyl.

- 20 6. The LPC of claim 5, wherein: X^1 is OH, SH, NH_2 , COR^5 or $COOR^4$, where R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R^5 is halide, heteroaryl or pseudohalide.

- 25 7. The LPC of claim 5, wherein Z is a group with three or more points of attachment: one to A, E, or the cyclic nucleus, and the others to two or more X^1 groups.

8. The LPC of claim 1, wherein the LPC has any of formulae:

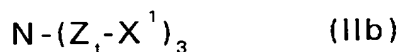
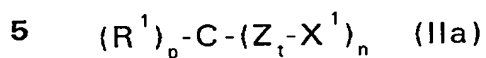
30



where in x is 0-6.

9. The LPC of claim 5, wherein: A is carbon and E is nitrogen.

10. The LPC of claim 5, wherein the LPC has formulae (IIa) or (IIb):



10 11. The LPC of claim 10, wherein p is 0 and n is 4.

12. The LPC of claim 11, wherein Z is any combination of 1-12 units selected from 1,4-phenylene and methylene, which units may be combined in any order.

13. The LPC of claim 10, wherein Z is C_{1-12} alkylene.

15 14. The LPC of claim 10, wherein X^1 is OH, SH or NH_2 .

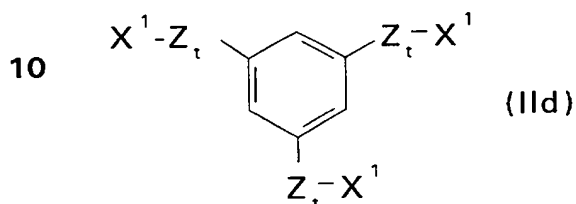
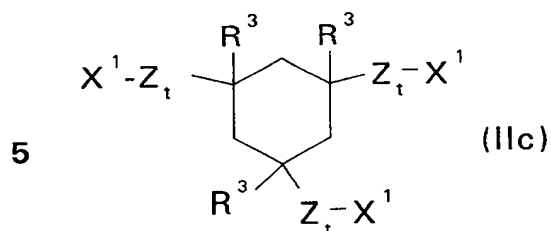
15. The LPC of claim 14, wherein X^1 is OH.

16. The LPC of claim 14, wherein X^1 is NH_2 .

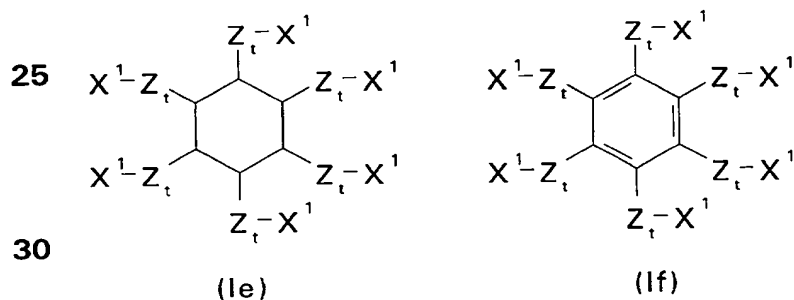
17. The LPC of claim 5, wherein the LPC has formulae (IIc) or (IId):

20

25



- 15
18. The LPC of claim 17, wherein Z is any combination of 1-12 units selected from 1,4-phenylene and methylene, which units may be combined in any order.
19. The LPC of claim 17, wherein Z is C₁₋₁₂ alkylene.
- 20
20. The LPC of claim 17, wherein X¹ is COR⁵ or COOR⁴.
21. The LPC of claim 17, wherein X¹ is COOR⁴.
22. The LPC of claim 5, wherein the LPC has formulae (Ie) or (If):



23. The LPC of claim 22, wherein Z is any combination of 1-12 units selected from 1,4-phenylene and methylene, which units may be combined in any order.
24. The LPC of claim 22, wherein Z is C₁₋₁₂ alkylene.
25. The LPC of claim 22, wherein X¹ is COR⁵ or COOR⁴.
26. The LPC of claim 25, wherein X¹ is COOR⁴.

27. The LPC of claim 1, wherein the LPC has formula $\text{Sp}(\text{O}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}_2)_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}_2)_n$, $\text{Sp}(\text{O}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}_2)_n$, $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{O}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{O}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{O}-(\text{CH}_2)_x-\text{SH})_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{O}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{S}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{S}-(\text{CH}_2)_x-\text{SH})_n$, $\text{Sp}(\text{O}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{S}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{O}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{S}-(\text{CH}_2)_x-\text{SH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{CO}-\text{O}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{CO}-\text{O}-(\text{CH}_2)_x-\text{SH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{CO}-\text{S}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{CO}-\text{S}-(\text{CH}_2)_x-\text{SH})_n$, $\text{Sp}(\text{C}(\text{O})-\text{O}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{C}(\text{O})-\text{S}-(\text{CH}_2)_x-\text{OH})_n$, $\text{Sp}(\text{C}(\text{O})-\text{O}-(\text{CH}_2)_x-\text{SH})_n$ or $\text{Sp}(\text{C}(\text{O})-\text{S}-(\text{CH}_2)_x-\text{SH})_n$ where x is 0-6.

28. The LPC of claim 27, wherein x is 2.

29. The LPC of claim 1 that is coupled to a photocleavable linker.

30. The LPC of claim 1 selected from the group consisting of $\text{Sp}(\text{O}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{S}-(\text{CH}_2)_2-\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, $\text{Sp}(\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$ and $\text{Sp}(\text{C}(\text{O})-\text{NH}-(\text{CH}_2)_x-\text{NH}-\text{C}(\text{O})-(\text{CH}_2)_x-\text{COOH})_n$, where x is 0-6.

31. The LPC of claim 1, selected from the group consisting of tetrakis(8-amino-6-aza-2-oxa-5-oxooctyl)methane, tetrakis(11-carboxy-6,9-diaza-5,10-dioxo-2-oxaundecyl)methane, tris(3-aza-6-carboxy-4-oxohexyl)amine, 1,3,5-benzenetricarboxylic acid tris-N-(2-aminoethyl)amide, 1,3,5-benzenetricarboxylic acid tris-N-(3-aza-6-carboxy-4-oxohexyl)amide, tetrakis{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-dT)₄-PE-LPC), 1,3,5-tris{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenyl-methyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC), tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-

trioxononyl]-benzene (dT₃-Aryl-LPC), tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine ((DMT-dT)₃-Amine-LPC) and tris[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

- 5 32. The LPC of claim 1 selected from the group consisting of tetrakis(11-carboxy-6,9-diaza-5,10-dioxo-2-oxaundecyl)methane, tris(3-aza-6-carboxy-4-oxohexyl)amine, 1,3,5-benzenetricarboxylic acid tris-N-(3-aza-6-carboxy-4-oxohexyl)amide, tetrakis{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-2-oxa-5,10,13-
- 10 trioxotridecyl}methane ((DMT-dT)₄-PE-LPC), 1,3,5-tris{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenyl-methyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC), tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-
- 15 trioxononyl]-benzene (dT₃-Aryl-LPC), tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine ((DMT-dT)₃-Amine-LPC) and tris[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

- 20 33. A method of solution phase biopolymer synthesis, comprising the steps of:
- (a) reacting an LPC of formula Sp(X¹)_n with a first monomer N¹;
- (b) separating and purifying the product of step (a) to afford a compound of formula Sp(X¹-N¹)_n;
- (c) reacting the product of step (b) with a second monomer N², a
- 25 dimer N²-N³ or a trimer N²-N³-N⁴; and
- (d) repeating steps (b) and (c) to produce an LPC-bound biopolymer of formula Sp(X¹-N¹-N²-...-N^m)_n, where m is 3 to 100, wherein:
- Sp is a polyvalent group that has more than two points of attachment, n corresponds to the number of points of attachment in Sp
- 30 and X¹ is a reactive group for biopolymer synthesis;
- N¹, N², N³...N^m are biopolymer monomers; and

the dimers and trimers comprise the monomers.

34. The method of claim 33, wherein the biopolymer is an oligonucleotide, peptide, peptide nucleic acid (PNA) or oligosaccharide.

35. The method of claim 33, further comprising the step of:

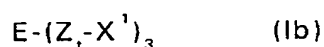
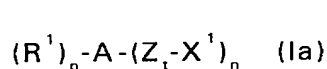
5 (e) cleaving the biopolymer from the LPC.

36. The method of claim 33, wherein the biopolymer is an oligonucleotide.

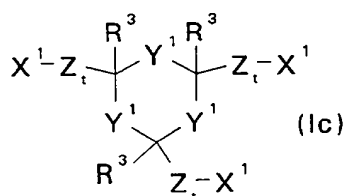
37. The method of claim 33, wherein n is 3-6.

38. The method of claim 33, wherein the LPC has formulae (I):

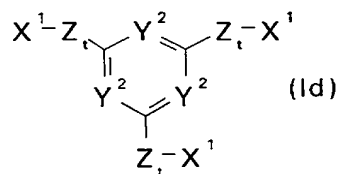
10



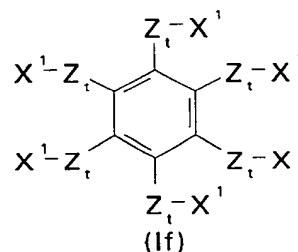
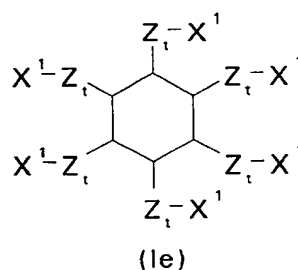
15



20



25



wherein: A is carbon or silicon; E is nitrogen or P(O); R¹ and R³ are each independently hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl,

30 heterocyclyl or heterocyclylalkyl; p is 0 or 1; Z is any combination of 0-12 units selected from 1,2-, 1,3- or 1,4-phenylene and alkylene, which units may be combined in any order; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis; n is 3 or 4; Y¹ is CH₂, NH, S or O; Y² is selected from CH and N; R¹, R³, X¹, Y¹, Y² and Z are unsubstituted
35 or substituted with one or more substituents each independently selected from Q; and Q is halogen, hydroxy, nitrile, nitro, formyl, mercapto, carboxy, alkyl, haloalkyl, polyhaloalkyl, aminoalkyl, diaminoalkyl, alkenyl

- containing 1 to 2 double bonds, alkynyl containing 1 to 2 triple bonds, cycloalkyl, cycloalkylalkyl, aryl, heteroaryl, arylalkyl, heteroarylalkyl, alkylidene, arylalkylidene, alkylcarbonyl, arylcarbonyl, heteroarylcarbonyl, alkoxycarbonyl, alkoxycarbonylalkyl, aryloxy, aryloxyalkyl, aryloxyalkylalkyl, aminocarbonyl, alkylaminocarbonyl, dialkylaminocarbonyl, arylamino-
- 5 carbonyl, diarylaminocarbonyl, arylalkylaminocarbonyl, alkoxy, aryloxy, perfluoroalkoxy, alkenyloxy, alkynyloxy, arylalkoxy, amino, aminoalkyl, alkylaminoalkyl, dialkylaminoalkyl, arylaminoalkyl, diarylaminoalkyl, alkylamino, dialkylamino, arylamino, diarylamino, alkylaryl amino, alkylcar-
- 10 bonylamino, alkoxycarbonylamino, arylcarbonylamino, aryloxy carbonyl-amino, azido, alkylthio, arylthio, perfluoroalkylthio, thiocyno, isothio-
- cyno, alkylsulfinyl, alkylsulfonyl, arylsulfinyl, arylsulfonyl, aminosulfonyl, alkylaminosulfonyl, dialkylaminosulfonyl, arylaminosulfonyl or diarylamino-
- sulfonyl.
- 15 39. The method of claim 38, wherein X^1 is OH, SH, NH_2 , COR^5 or $COOR^4$, where R^4 is selected from hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl and heterocyclylalkyl; and R^5 is halide, heteroaryl or pseudohalide.
40. The method of claim 33, wherein the LPC is selected from the
- 20 group consisting of tetrakis(11-carboxy-6,9-diaza-5,10-dioxo-2-oxaundecyl)methane, tris(3-aza-6-carboxy-4-oxohexyl)amine, 1,3,5-benzenetricarboxylic acid tris-N-(3-aza-6-carboxy-4-oxohexyl)amide, tetrakis{6,9-diaza-13-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-
- 25 deoxythymidine-3'-O-yl]-2-oxa-5,10,13-trioxotridecyl}methane ((DMT-dT)₄-PE-LPC), 1,3,5-tris{2,5-diaza-9-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-1,6,9-trioxononyl}-benzene ((DMT-dT)₃-Aryl-LPC), tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-
- 5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-
- 30 LPC), tris-{3-aza-4,7-dioxo-7-[5'-O-(4,4'-dimethoxytriphenylmethyl)-2'-deoxythymidine-3'-O-yl]-heptyl}-amine ((DMT-dT)₃-Amine-LPC) and tris[3-

aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

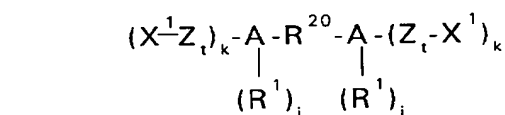
41. The method of claim 33, wherein the LPC is selected from tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC), and tris[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

42. The method of claim 33, wherein the LPC is 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC).

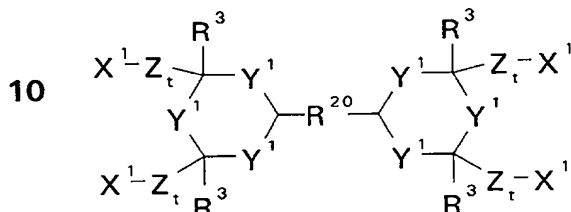
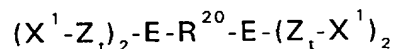
43. The LPC of claim 1 selected from the group consisting of tetrakis[13-(2'-deoxythymidin-3'-O-yl)-6,9-diaza-2-oxa-5,10,13-trioxotridecyl]-methane (dT₄-PE-LPC), 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC), and tris[3-aza-7-(2'-deoxythymidine-3'-O-yl)-4,7-dioxoheptyl]-amine (dT₃-Amine-LPC).

44. The LPC of claim 1 that is 1,3,5-tris[9-(2'-deoxythymidin-3'-O-yl)-2,5-diaza-1,6,9-trioxononyl]-benzene (dT₃-Aryl-LPC).

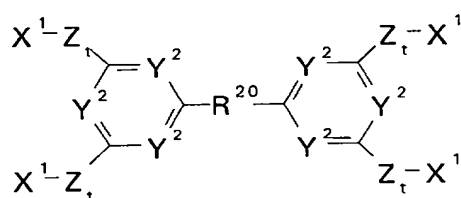
45. The LPC of claim 1 that has formulae:



5



15

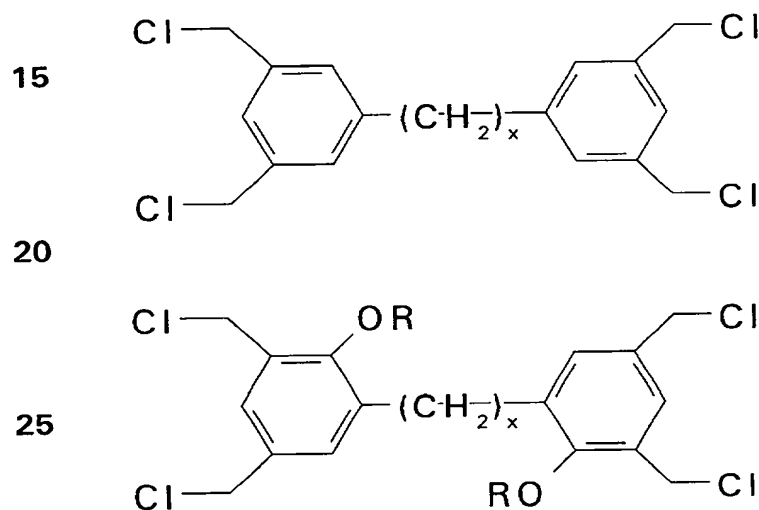


20

wherein: A is carbon or silicon; E is nitrogen or P(O); R¹ and R³ are each independently hydrogen, alkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocyclyl or heterocyclylalkyl; Z is any combination of 1-12 units selected from 1,2-, 1,3- or 1,4-phenylene and alkylene, which units may be combined in any order, with the proviso that if the LPC is of formula (Ia) or (Ib), then Z contains at least two phenylene or methylene units; t is 0 or 1; X¹ is any reactive group which can be used in biopolymer synthesis; Y¹ is CH₂, NH, S or O; Y² is selected from CH and N; R¹, R³, X¹, Y¹, Y² and Z are unsubstituted or substituted with one or more substituents each independently selected from Q; and Q is halogen, hydroxy, nitrile, nitro, formyl, mercapto, carboxy, alkyl, haloalkyl, polyhaloalkyl, aminoalkyl, diaminoalkyl, alkenyl containing 1 to 2 double bonds, alkynyl containing 1 to 2 triple bonds, cycloalkyl, cycloalkylalkyl, aryl, heteroaryl, arylalkyl, heteroarylalkyl, alkylidene, arylalkylidene, alkylcarbonyl, arylcarbonyl, heteroarylcarbonyl, alkoxy carbonyl, alkoxy carbonylalkyl, aryloxy carbonyl, aryloxy carbonylalkyl, aminocarbonyl, alkyl-

- aminocarbonyl, dialkylaminocarbonyl, arylaminocarbonyl, diarylamino-
carbonyl, arylalkylaminocarbonyl, alkoxy, aryloxy, perfluoroalkoxy, alkenyl-
oxy, alkynyloxy, arylalkoxy, amino, aminoalkyl, alkylaminoalkyl, dialkyl-
aminoalkyl, arylaminoalkyl, diarylaminoalkyl, alkylamino, dialkylamino,
5 arylamino, diarylamino, alkylaryl amino, alkylcarbonylamino, alkoxycarbon-
ylamino, arylcarbonylamino, aryloxycarbonylamino, azido, alkylthio, aryl-
thio, perfluoroalkylthio, thiocyano, isothiocyano, alkylsulfinyl, alkyl-
sulfonyl, arylsulfinyl, arylsulfonyl, aminosulfonyl, alkylaminosulfonyl,
dialkylaminosulfonyl, arylaminosulfonyl or diarylaminosulfonyl; R^{20} is
10 alkylene, alkenylene, alkynylene, arylene or heteroarylene; k is 2 or 3; and
 j is 0 or 1.

46. The LPC of claim 45, wherein the compound has the formulae:



- 30 wherein x is 0-6 and R is lower alkyl.

47. The method of claim 33, wherein the monomers are nucleotides, nucleosides, natural or unnatural amino acids, protein nucleic acid (PNA) monomers or monosaccharides.

48. A method of solution phase biopolymer synthesis, comprising
35 the steps of:

(a) reacting an LPC of formula $Sp(X^1)_n$ with a first monomer N^1 ;

(b) separating and purifying the product of step (a) to afford a compound of formula $\text{Sp}(\text{X}^1\text{-N}^1)_n$;

(c) reacting the product of step (b) with a second monomer N^2 , a dimer $\text{N}^2\text{-N}^3$ or a trimer $\text{N}^2\text{-N}^3\text{-N}^4$; and

- 5 (d) repeating steps (b) and (c) to produce an LPC-bound biopolymer of formula $\text{Sp}(\text{X}^1\text{-N}^1\text{-N}^2\text{-...-N}^m)_n$, where m is 3 to 100, wherein:

Sp is a polyvalent group that has two or more points of attachment, n corresponds to the number of points of attachment in Sp and X^1 is a reactive group for biopolymer synthesis;

- 10 N^1 , N^2 , $\text{N}^3\text{...N}^m$ are biopolymer monomers;
the dimers and trimers comprise the monomers; and
the protocol used in steps (c) and (d) to synthesize the biopolymer, preferably the oligonucleotide, is the phosphoramidite protocol.

49. The LPC of claim 1 coupled to a biopolymer.

ABSTRACT

Multifunctional liquid phase carriers (LPCs) and methods of using LPCs for the preparation of biopolymers are provided. The LPCs are highly symmetrical compounds that possess more than two points of attachment for biopolymer synthesis. The LPCs have the formula $Sp(X^1)_n$, where Sp is a highly symmetrical moiety such that all X^1 groups are equivalent. X^1 is a functional group that is suitable for biopolymer synthesis, including OH, SH, NH_2 , COOH and the like. Biopolymers that may be produced using the methods provided include oligonucleotides, peptides, protein nucleic acids (PNAs) and oligosaccharides. Analogs of the biopolymers may also be prepared using the methods.

DECLARATION FOR PATENT APPLICATION

As below-named inventors, we hereby declare that:

Our residences, post office addresses and citizenships are as stated below next to our names.

We believe we are the original, first and joint inventors of the subject matter which is claimed and for which a patent is sought on the invention entitled:

SOLUTION PHASE BIOPOLYMER SYNTHESIS

the specification of which

- (X) is attached hereto.
 () was filed by an authorized person on my behalf on April 27, 1998 as Application Serial No. _____.

We hereby state that we have reviewed and understand the contents of the above-identified specification, including the claims as amended by any amendment referred to above.

We acknowledge the duty to disclose information which is material to the examination of this application in accordance with Title 37, Code of Federal Regulations, §1.56(a).

We hereby claim foreign priority benefits under Title 35, United States Code, §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate listed below and so identified, or §365(a) of any PCT international application that designated at least one country other than the United States of America, listed below, and we have also identified below any foreign application for patent or inventor's certificate or PCT international application on this invention filed by us or our legal representatives or assigns and having a filing date before that of the application on which priority is claimed.

<u>Number</u>	<u>Country</u>	<u>Day/Month/Year Filed</u>	<u>Priority Claimed (Yes or No)</u>
N/A			

We hereby claim benefit under Title 35, United States Code, §119(e) of any United States provisional application(s) listed below:

<u>Application Serial No.</u>	<u>Filing Date</u>
N/A	

We hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States application in the manner provided by the first paragraph of Title 35, United States Code, §112, we acknowledge the duty to disclose material information as defined in Title 37, Code of Federal Regulations, §1.56(a) which occurred between the filing date of the prior application and the national or PCT international filing date of this application:

<u>Application Serial No.</u>	<u>Filing Date</u>	<u>Status</u>
-------------------------------	--------------------	---------------

N/A

<u>PCT Application No.</u>	<u>Filing Date</u>	<u>Status</u>
----------------------------	--------------------	---------------

N/A

We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

We hereby appoint the following attorneys and agents, with full power of substitution and revocation, to prosecute this application and to transact all business in the United States Patent and Trademark Office connected therewith and request that all correspondence and telephone calls in respect to this application be directed to Stephanie Seidman, BROWN, MARTIN, HALLER & McCLAIN, 1660 Union Street, San Diego, California 92101-2926:

<u>Attorney</u>	<u>Reg. No.</u>
Stephanie Seidman	33,779
Paula K. Schoeneck	39,362
Carl Brown	18,587
Neil R. Martin	23,088
John L. Haller	27,795
James W. McClain	24,536
Eleanor Musick	35,623

Address for correspondence: Stephanie Seidman
BROWN, MARTIN, HALLER & McCLAIN
1660 Union Street
San Diego, CA 92101-2926

Full name of joint inventor: Hubert Köster

Inventor's signature: _____

Date: _____

Residence: La Jolla, California

Post Office Address: 8636C Via Mallorca Drive,
La Jolla, California 92037, United States of America

Citizenship: Germany

Full name of joint inventor: Ralf Wörl

Inventor's signature: _____

Date: _____

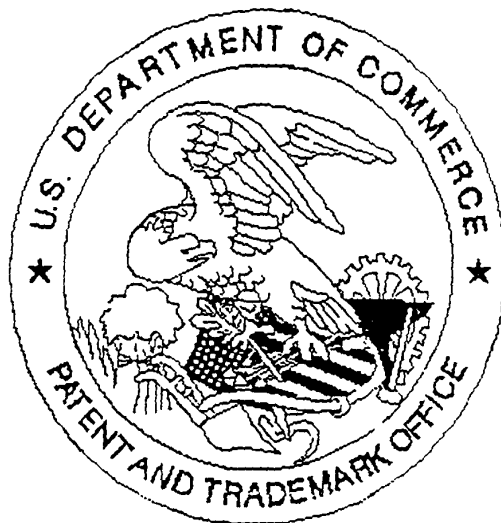
Residence: _____

Post Office Address: _____

Citizenship: Germany

United States Patent & Trademark Office

Office of Initial Patent Examination -- Scanning Division



Application deficiencies found during scanning:

- 1 Application papers are not suitable for scanning and are not in compliance with 37 CFR 1.52 because
 - ☐ All sheets must be the same size and either A4 (21 cm x 29.7 cm) or 8-1/2" x 11"
 - Pages _____ do not meet these requirements
 - ☐ Papers are not flexible, strong, smooth, non-shiny, durable, and white.
 - ☐ Papers are not typewritten or mechanically printed in permanent ink on one side.
 - ☐ Papers contain improper margins. Each sheet must have a left margin of at least 2.5 cm (1") and top, bottom and right margins of at least 2.0 cm (3/4").
 - ☐ Papers contain hand lettering.
- 2 Drawings are not in compliance and were not scanned because.
 - ☐ The drawings or copy of drawings are not suitable for electronic reproduction.
 - ☐ All drawings sheets are not the same size. Pages must be either A4 (21 cm x 29.7 cm) or 8-1/2" x 11"
 - ☐ Each sheet must include a top and left margin of at least 2.5 cm (1"), a right margin of at least 1.5 cm (9/16") and a bottom margin of at least 1.0 cm (3/8")
- 3 Page(s) _____ are not of sufficient clarity, contrast and quality for electronic reproduction.
- 4 Page(s) _____ are missing.
- 5 OTHER NO DRAWINGS